## **Final Scoping Memorandum**

### **Bremerton Gas Works Site**

Prepared for: Cascade Natural Gas Corporation

Aspect Project No. 080239-005 •Anchor QEA Project No. 131014-01.01

March 5, 2015

Prepared by



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#### **Acronyms**

Anchor QEA Anchor QEA, LLC

AOC Administrative Settlement Agreement and Order on

Consent for Remedial Investigation Feasibility Study

ARAR applicable or relevant and appropriate requirement

Aspect Consulting, LLC

AST aboveground storage tank

BTAG Biological Technical Assistance Group

BTEX benzene, toluene, ethylbenzene, and xylenes

Cascade Natural Gas Corporation

CERCLA Comprehensive Environmental Response,

Compensation, and Liability Act

CFR Code of Federal Regulations

City City of Bremerton

COC contaminant of concern

COPC contaminant of potential concern

cPAH carcinogenic polycyclic aromatic hydrocarbon

CSM conceptual site model

CSO combined sewer overflow

DNAPL dense non-aqueous phase liquid

DNR Washington State Department of Natural Resources

DQO data quality objective

DU data usability

E&E Ecology & Environment

Ecology Washington State Department of Ecology

EcoSSL ecological soil screening level

ENVVEST Environmental Investment Project

EPA U.S. Environmental Protection Agency

ERA ecological risk assessment

ER-L effect range-low

ER-M effect range-medium

FS Feasibility Study

HHRA human health risk assessment

HPAH high-molecular-weight polycyclic aromatic

hydrocarbon

ISA Initial Study Area

KPHD Kitsap Public Health District

Lent's Lents and Blombergs

LNAPL light non-aqueous phase liquid

LPAH low-molecular-weight polycyclic aromatic

hydrocarbon

MCL maximum contaminant level

μg/kg microgram(s) per kilogram

μg/L microgram(s) per liter

MDAC minimum data acceptability criteria

mg/kg milligram(s) per kilogram

MGP manufactured gas plant

MTCA Washington State Model Toxics Control Act

NAPL non-aqueous phase liquid

NOAA National Oceanographic and Atmospheric

Administration

NPL National Priorities List

NRHP National Register of Historic Places

Order Administrative Order for a Pollution Incident

PAH polycyclic aromatic hydrocarbon

PCB polychlorinated biphenyl

PCP pentachlorophenol

PHS Priority Habitats and Species Program

PQL practical quantitation limit

PRG preliminary remediation goal

PSAMP Puget Sound Assessment and Monitoring Program

RAO remedial action objectives

RCRA Resource Conservation and Recovery Act

RI Remedial Investigation

RSL regional screening level

SCO sediment cleanup objective

Site Bremerton Gas Works Site

SMS Sediment Management Standards

SOW Statement of Work

SVOC semivolatile organic compound

TBA Targeted Brownfields Assessment

TBC to be considered

TCRA time critical removal action

TEQ toxic equivalent

TOC total organic carbon

TPH total petroleum hydrocarbons

Tribe Suquamish Tribe

TS total solids

USCG U.S. Coast Guard

UST underground storage tank

VOC volatile organic compound

WDFW Washington Department of Fish and Wildlife

### 1 Introduction

Cascade Natural Gas Corporation (Cascade) is conducting a Remedial Investigation (RI) and Feasibility Study (FS) at the Bremerton Gas Works Site (Site) in Bremerton, Washington, under the direction of the U.S. Environmental Protection Agency (EPA). The work is being conducted in accordance with the Administrative Settlement Agreement and Order on Consent for Remedial Investigation Feasibility Study (AOC; Comprehensive Environmental Response, Compensation, and Liability Act [CERCLA] Docket No. 10-2013-0104). Consistent with the AOC, the Site includes the area where the gas works was formerly located (Figure 1-1), the adjacent beach, and the associated areal extent of contamination.

In November 2010, Cascade performed a time critical removal action (TCRA) at the Site with oversight from the EPA and the U.S. Coast Guard (USCG). EPA placed the Site on the National Priorities List (NPL) on May 10, 2012, and the AOC was executed on May 1, 2013. In accordance with the AOC, a Removal Evaluation and a Removal Action were performed in 2013 to assess and mitigate potential threats to human health, human welfare, and the environment attributable to site-related contaminants prior to completion of the RI/FS. That work is documented in the Removal Evaluation Report (Anchor QEA and Aspect Consulting 2013c) and the Time-Critical Removal Action Report (Anchor QEA and Aspect Consulting 2014).

Scoping is the initial planning phase of the RI/FS, as described in the Guidance for Conducting Remedial Investigations and Feasibility Studies under CERCLA (EPA 1988a). In accordance with the AOC, this Scoping Memorandum summarizes existing information to identify the scope of data collection needed to complete the RI/FS. After completion of the Scoping Memorandum, detailed project planning, including proposal of specific work to address data gaps, will be conducted and documented in the RI/FS Work Plan.

The objectives of the scoping process and the content and organization of this Scoping Memorandum are described below.

### 1.1 Scoping Objectives

The goal of scoping is to present and evaluate known information to identify the scope of data/information gathering necessary to conduct the RI and FS for the Site. Specific objectives of the scoping process are as follows:

- Identify and compile applicable historical information and data that are of acceptable quality for use during the RI/FS process;
- Identify relevant existing studies regarding the characteristics of environmental media and the condition of receptor populations at the Site;

- Identify usable information and data from current and historical studies for use in developing a conceptual site model (CSM);
- Identify an Initial Study Area (ISA) for both the upland area and the sediment area of the Site;
- Identify Site-specific objectives of the RI/FS, including initial preliminary remediation goals (PRGs), to help evaluate the adequacy of the existing information and to identify any data gaps;
- Establish a preliminary list of applicable and relevant and appropriate requirements (ARARs);
- Identify potential remedial approaches or technologies that may be applied, to determine potential data needs associated with remedial alternative development; and
- Document the need for additional information and data to the extent practicable to support the RI/FS.

### 1.2 Document Organization

The remainder of this Scoping Memorandum is organized as follows:

- Section 2 Site History and Description describes the Site location, ownership, zoning, and operational history.
- Section 3 Environmental Setting describes the Site physical conditions including topography and surface drainage, geology and hydrogeology, ecological environment (terrestrial and aquatic), cultural resources, land use, and existing infrastructure.
- Section 4 Previous Investigations and Remedial Actions describes the previous investigations and cleanup actions conducted at the Site and on surrounding properties.
- Section 5 Preliminary Conceptual Site Model identifies the potential sources
  of contaminants, their potential migration pathways, the environmental media in
  which their presence is suspected or has been confirmed, and the potential
  contaminant exposure pathways and receptors.
- **Section 6 Project Planning** identifies potential ARARs for the Site, initial PRGs for potential contaminants in environmental media, and preliminary remedial action objectives (RAOs).
- Section 7 Existing Data and Data Usability summarizes data collected during previous Site investigations and evaluates the quality and usability of that data.

- Section 8 RI/FS Approach identifies preliminary contaminants of potential concern (COPCs), defines the ISA to be investigated, provides an overview of the risk assessment approach, and identifies potential remedial approaches.
- Section 9 Summary and Data Gaps compiles the principal data needs for the RI/FS as defined in this Scoping Memorandum and defines the anticipated sequence of investigation activities.
- **Section 10 References** lists documents used as sources of information and referenced in this Scoping Memorandum.

### 2 Site History and Description

This section describes the property upon which the former gas works was located and the properties surrounding the former gas works and discusses the operational and regulatory history of those properties.

### 2.1 Site Location and Description

The former gas works was located between Thompson Drive and Pennsylvania Avenue (Figure 2-1) on approximately 2.8 acres of property along the south shore of Port Washington Narrows in Bremerton, Washington. The historical street addresses for the former gas works included 1720 and 1800 Thompson Drive.

The real property upon which the former gas works was located (Former Gas Works Property) relative to current parcel boundaries is shown on Figure 2-1. Due to a boundary line adjustment in 1992, the Former Gas Works Property includes portions of two existing tax parcels:

- Kitsap County Parcel No. 3711-000-0010-0409 (McConkey Property). This parcel
  is owned by the McConkey Family Trust. The former gas works covered the entire
  parcel. No current or historical street address has been identified for this parcel.
- Kitsap County Parcel No. 3741-000-022-0101 1701 Pennsylvania Avenue (Sesko Property). This parcel is owned by Natasha Sesko. The former gas works covered the northwestern portion of this parcel.

The following properties are located near the Former Gas Works Property and have had either suspected or confirmed releases of contaminants from historical operations unrelated to the former gas works:

- 1723 Pennsylvania Avenue (Penn Plaza Property). This property is owned by Penn Plaza Storage, LLC. There are multiple street addresses associated with this property, but it is listed in the Kitsap County assessor's database as 1723 Pennsylvania Avenue.
- 1701 Thompson Drive (Former ARCO Property). This property is owned by Pipeworks Mechanical & Service, Inc. It is located southwest of the Former Gas Works Property, across Thompson Drive.
- 1702 Pennsylvania Avenue (Former SC Fuels Property). This property is owned by NFS Properties 2, LLC. It is located east of the Sesko Property, across Pennsylvania Avenue.

The Port Washington Narrows is located north of the McConkey, Sesko, and Former SC Fuels Properties. The Port Washington Narrows consists of aquatic lands owned by the

State of Washington and managed by the Washington State Department of Natural Resources (DNR).

#### 2.2 Site Uses Prior to 1930

The Port Washington Narrows and the adjacent uplands are located in the traditional territory of the Suquamish Tribe (Tribe), a Southern Coast Salish community speaking a dialect of the Southern Lushootseed language (Suttles and Lane 1990). Shoreline locations in Dyes Inlet would have been available after stabilization of sea levels in the mid-Holocene (Thorson 1980); therefore, Native American use of the area may date back more than 5,000 years. A variety of traditional activities took place in the general vicinity. In 1855, the Tribe signed the Treaty of Point Elliott, which ceded lands and established the reservation at Port Madison. The Tribe retained "the right of taking fish at usual and accustomed grounds and stations" (Treaty of Point Elliott 1855), and the Port Washington Narrows is within the Tribe's adjudicated Usual and Accustomed area.

### 2.3 Current and Historical Use and Operations

Historical use and operations on the properties and aquatic lands are based on historical records, including aerial photographs, interviews with current and former workers, owners, area residents, historical maps, deeds, Washington State Department of Ecology (Ecology) records, City of Bremerton (City) records, and DNR lease records. A number of historical documents are included in previous assessments of historical Site use (TechLaw 2006; Hart Crowser 2007). Available and relevant historical records are provided in Appendix A for reference.

Historical and current operations on the Former Gas Works Property (which consists of the entire McConkey Property and a portion of the Sesko Property) as well as historical and current operations on the other portion of the Sesko Property are described in Section 2.3.1. Historical and current operations on adjoining properties are described in Section 2.3.2.

### 2.3.1 Operations on McConkey and Sesko Properties

#### 2.3.1.1 Former Gas Works Operations

In 1930, the Former Gas Works Property was developed as a gas works (a.k.a., manufactured gas plant, or MGP). Gas works were a common industry in large and small towns throughout the United States and Europe from approximately the mid-1800s to the mid-1900s. At a gas works, coal, coke, and/or petroleum products were heated in furnaces to produce manufactured gas, which was subsequently distributed via a gas piping network to the surrounding homes and businesses for heating, cooking, and lighting. Gas works used or generated a number of products and byproducts, including non-aqueous phase liquids (NAPLs) such as oils and tars, aqueous waste streams, and solid materials containing chemicals that may pose a risk to human health or the

environment because they are toxic or carcinogenic (resulting in cancer effects). These contaminants include hydrocarbons such as benzene, toluene, ethylbenzene, and xylenes (BTEX) and polycyclic aromatic hydrocarbons (PAHs), which can persist for a long time in the environment. Contaminant releases from historical gas works operations at other locations have resulted in sites where contamination remains in the subsurface as NAPLs, sorbed to soil or sediments or dissolved in the groundwater.

Because of the potential hazards posed by historical gas works facilities, these facilities are often the focus of state-led or federally led efforts to investigate and clean up contamination to protect human health and the environment. To characterize and remediate these facilities, it is important to understand traditional gas works operations, the types of contaminants that may be present, and where contaminants may have been released. This section provides a summary of what is known about operations at the former gas works based on historical documentation and what is assumed based on typical gas works operations. This section also identifies the contaminants usually associated with gas works feedstocks, fuels, and byproducts that may be present at the Site. Uncertainties about historical practices and potential releases will be addressed in the RI though field investigations. Further discussion of potential release mechanisms and transport of contaminants in the subsurface is provided in Section 5, Preliminary Conceptual Site Model.

The operational history of the former gas works is as follows:

- **1930 to 1931.** The former gas works was constructed by the Western Gas and Utilities Corporation. It included a dock on aquatic lands initially leased from DNR on November 25, 1930 (Former Gas Works Dock).
- 1931 to 1955. Manufactured gas was produced using the carbureted water-gas process, from feedstocks of coal, coke briquettes, and petroleum products. In the 1940s, a standby plant for producing natural gas by blending liquefied petroleum (butane or propane) and air was installed. Gas produced at the Former Gas Work Property in the 1940s and 1950s was from manufactured gas and from butane-air. In approximately 1955 (Simonson 1997b), manufactured gas operations ceased, and all gas was produced from butane-air mixing.

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<sup>&</sup>lt;sup>1</sup> In 1931, the Western Gas and Utilities Corporation changed its name to the Western Gas Company of Washington. The Western Gas and Utilities Corporation and the Western Gas Company of Washington are collectively referred to as "Western" herein.

<sup>&</sup>lt;sup>2</sup> Typically, diesel-range fuel oils were used for petroleum feedstock for the carbureted water-gas process (Hatheway 2012). However, one historical map (Sanborn 1946) indicates gasoline and fuel oil were stored in the northeast corner of the Former Gas Works Property.

- **1955 to 1963.** Natural gas was produced from butane-air mixing. In 1963, with the completion of a natural gas pipeline to the region, gas production ceased.
- 1963 to 1972. Some of the structures and tanks were removed between 1964 and 1965, and the concrete piers supporting the tanks were jackhammered and hauled away (White 1998). The former plant building was reportedly used for pipe storage and, for a short time, magnesium mining research (*Bremerton Sun* 1972). In 1972, the remaining structures, including the former plant building, were sold and dismantled.

In 1972, the Former Gas Works Property was acquired by Harold D. and L. Irene Lent and Theodore and Marian J. Blomberg, doing business as "Lent, Blomberg, Lent." The Lent and Blomberg families operated several businesses in the vicinity of the Former Gas Works Property, including an oil distribution business on the Sesko Property under the name Lents, Inc. (see further discussion in Section 2.3.1.3). All entities and individuals associated with the Lents and Blombergs are referred to in this Scoping Memorandum as "Lent's."

In 1979, Paul and Margaret McConkey acquired the majority of the Former Gas Works Property. The McConkeys acquired the remainder of the Former Gas Works Property in 1985. A portion of the Former Gas Works Property was sold to William Sesko in 1992.

The summary of gas works operations provided in this section combines available historical information about the layout and operations of the former gas works with information compiled from multiple sources regarding the operations of typical manufactured gas facilities, including generated byproducts and likely sources of releases of hazardous substances. Whereas this summary provides an overview of operations at the former gas works, it likely does not provide a complete picture of all sources, disposal areas, and spills and/or releases that may have occurred, which will be investigated primarily through the collection and evaluation of data during the RI. Chemical feedstocks and potential byproducts typical of carbureted water-gas production<sup>3</sup> include the following:

•	Feedstock and Fuels: Gasoline, Diesel, Coal, or Coke Briquettes. The
	contaminants potentially associated with feedstock and fuels include the
	following:

0	BTEX;	

<sup>&</sup>lt;sup>3</sup> Two byproducts typically generated at coal and/or oil gas plants, ammoniacal liquor and lampblack (carbon soot), were generally not generated in significant quantities by the carbureted water-gas process (Hatheway 2012).

- Naphthalenes; and
- o PAHs.
- Byproducts: Light Oil, Carbureted Water-Gas Tar, Ash, Clinker, Slag, Soot, and Spent Purifier Filter Media. The contaminants potentially associated with byproducts include the following:
  - O BTEX:
  - Naphthalenes;
  - o PAHs;
  - o Phenols; and
  - Other semivolatile organic compounds (SVOCs), including creosol, carbazole, and dibenzofuran.

Section 8.1 provides further discussion of the Site-specific COPCs.

Production of natural gas using liquefied petroleum (butane or propane) blended with air is not anticipated to have resulted in contamination of the subsurface because butane and propane are gases at atmospheric conditions.

A flow chart showing the gas works process as understood at the Site (based on available plant maps and typical carbureted water-gas operations), including the production of byproducts, is presented on Figure 2-2. The locations of key plant features are shown on Figure 2-3. The general sequence of operations is as follows:

- **Product Delivery and Storage.** Solid feedstocks (coal and coke briquettes) were transported to the Site by barge and offloaded via a winch to a storage slab located in the northwest corner of the Former Gas Works Property. Petroleum products were also delivered to the former gas works via barge and conveyed via a pipeline up the Former Gas Works Dock to storage tanks located in the northeast corner of the Former Gas Works Property.
- Gas Generation and Purification. These operations were located in the north-central portion of the Former Gas Works Property (Figure 2-3). Two generator sets (furnaces) were located in the main plant building: one in the northern portion of the building and one in the middle of the building (Simonson 1997b). The main plant building had a concrete floor (Simonson 1997b). Coal and coke were placed in the generators and heated, and fuel oil was sprayed into the generators to produce gas. The resulting gas stream was then passed through a series of devices to cool the gas and remove impurities. These devices are described below:

Scrubber. After gas generation comes clarification, in which tar is separated from the gas using a scrubber or similar equipment. These devices are typically located adjacent to the generator sets. A historical plant map shows the scrubber located directly west of the generator sets. A former plant worker indicated that the scrubber consisted of a tank with wooden slots and water to "wash out" the gas (Simonson 1997b). An engineer's report (Tymstra 1942) indicates that wood chips and excelsior (i.e., wood shavings) were used to remove tar from the gas.

The clarification process typically produced tar, tar-soaked wood chips or shavings, gas liquor (aqueous solutions containing dissolved and suspended tar particles), and tar-water emulsions. Light oils may also have been produced in the scrubbing process. Tar-water emulsions from scrubbers were typically removed from clarification equipment and transported to residual management areas to separate tar from the water (Hatheway 2012). The fate of byproducts and residuals is discussed in the bullet "Residuals Management."

- Gas Holder. A large gas holder was located south of the scrubber, west of the main plant building. The bottom of the gas holder was reportedly 15 feet deep and contained tar and water (Simonson 1997a). The materials used to construct the base of the gas holder are not known.
- o **Purifier.** Gas was passed through a bed of filter media to remove impurities such as sulfide from the gas. Typical filter media included wood chips and/or iron oxide. An engineer's report (Tymstra 1942) indicated that iron-oxide-covered chips were used at the gas works to remove sulfur compounds from gas. Multiple purifiers in parallel were typically installed to allow changeout of purifier media without interrupting the process (Hatheway 2012). Three purifiers were located at the Former Gas Works Property south of the large gas holder. In addition to the generation of spent purifier media, which included some accumulated tar (Tymstra 1942), some liquid streams (including tar, gas liquor, and light oil) may have condensed during purification and were typically manually removed from the purifier box (Hatheway 2012). The fate of these byproducts is discussed in the following bullet.
- Residuals Management. In addition to the gas produced by the manufactured gas process, residual materials were also produced and separated from the gas at several steps during the process. These residuals were intermediate waste streams typically managed on-site and further processed to create byproducts for disposal or reuse. Residuals from the manufactured gas process included the following:

**Tar-Water Emulsion.** Tar removed from the gas stream, particularly from the condenser, was often a tar-water emulsion. Tar required a low water content to be saleable. Tar-water emulsions were typically removed from clarification equipment and transported to residual management areas to separate the tar from the water (Hatheway 2012). Tar and water were typically separated by placing the emulsion in pits, cisterns, or tar wells (typically shallow boxes that may be lined or unlined) and allowing the tar to settle out. A former plant map shows tar wells and a residue cistern located west of the purifiers near the edge of the ravine adjacent to the former gas works (Former Ravine). A former resident recalled a tar pit located on the southwest corner of the Former Gas Works Property (Judd 2014), and an engineer's report (Tymstra 1942) noted, "The tar emulsion is dumped in shallow pits dug at random in the ground." A historical journal (Perry 2002) indicated that the former gas works "had a pond for dumping surplus creosote-type fluids. This would overflow and the material would go into the channel." It is unknown how tar-water emulsions were transported to these areas or how tar was transported from these areas to the tar storage tank, which was located on the south side of the Former Gas Works Property.

#### • Storage, Distribution, and Disposal of Gas and Byproducts.

- Finished Gas. Gas that had passed through the scrubbers and purifiers was pumped through compressors located in the engine room (south of the main plant building) and stored in finished gas storage tanks located south of the main operations area. Gas was piped from the finished gas tanks to the gas distribution system along an 8-inch-diameter gas main located in Thompson Avenue. Typically in manufactured gas distribution systems, a minor amount of oil would condense within the initial section of distribution piping, which would be collected in a drip tank located near the facility (Hatheway 2012). A drip tank located just south of the Former Gas Works Property (Figure 2-3) is shown on a historical plant sketch.
- Light Oil. Light oils typically contain one- or two-ring aromatic compounds, such as BTEX, and naphthalenes and have a density less than that of water (i.e., light, non-aqueous phase liquids [LNAPLs]). Light oils were sometimes reused in the carbureted water-gas process. According to a former worker, light oils were produced in small quantities at the former gas works and stored in a tank south of the finished gas storage tanks, and they were occasionally sprayed to control weeds in the southwest corner of the Former Gas Works Property or as automotive fuel for workers' vehicles (Simonson 1997b).

- Carbureted Water-Gas Tar. This tar typically contains both light aromatics (e.g., BTEX) and semivolatile hydrocarbons. Semivolatiles in coal tar primarily consist of PAHs but also include phenols and heterocyclic aromatics (i.e., carbazole or dibenzofuran). Coal tar is typically more dense than water (i.e., dense non-aqueous phase liquids [DNAPLs]). According to a former worker (Simonson 1997b), tar was a saleable product that was collected, stored in a tank on the south side of the Former Gas Works Property, and piped to barges at the Former Gas Works Dock. However, it is unlikely that all tar generated over the entire life span of the former gas works was recovered and sold in this manner.
- Gas Liquor. Gas liquor is water containing dissolved and suspended tar and oil constituents. According to the 1942 report (Tymstra 1942), this stream was discharged to "the bay" (i.e., the Port Washington Narrows) through a drainpipe.<sup>4</sup>
- Ash, Clinker, and Slag (Mineral Residue of Fuel and Feedstocks) from the Furnaces. Ash is generally powdery, whereas clinker is partially fused, and slag is fused. These materials were reportedly placed on the bluff along the shoreline (Judd 2014) north of the Former Gas Works Property and may have also been deposited in the Former Ravine.<sup>5</sup>
- Soot from the Furnaces. This material was reportedly placed in the Former Ravine near the oil storage tanks (Tymstra 1942).
- Spent Scrubber and Purifier Media. When scrubber and purifier media such as tar-soaked wood chips and shavings were saturated, they were removed and replaced. Spent scrubber media contains tar, and spent purifier media often contains tar, sulfide, and cyanide compounds removed during purification, including Prussian Blue (an iron-cyanide compound) (Hatheway 2012). During a period of gas works operations, tar-soaked wood chips and excelsior produced on-site were reportedly placed in the Former Ravine near the oil storage tanks (Tymstra 1942). However, an individual who worked at the former gas works between

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<sup>&</sup>lt;sup>4</sup> It is suspected that the drain pipe referred to in the 1942 report corresponds to the former outfall that was removed and plugged as part of the 2010 TCRA (see Section 4.2.1).

<sup>&</sup>lt;sup>5</sup> Boring logs for SP01 and MW04, which were located in the Former Ravine, indicate ash.

1953 and 1955 indicated that the spent purifier media were hauled off-site.

#### 2.3.1.2 Post-1972 Operations on the McConkey Property

Operations on the McConkey Property after the former gas works discontinued operations have included activities by Lent's between approximately 1972 and 1982 and industrial park operations by others from approximately 1982 to the present. Operations on the McConkey Property have included metal fabrication and sandblasting on the southern portion of the property and parking and equipment storage across the other portion of the property. Two buildings are located in the southern portion of the McConkey Property. Historical and current operations on the McConkey Property are shown on Figure 2-4. A generalized process flow diagram of the metal fabrication process is shown on Figure 2-5.

Ecology inspected industrial park operations on the McConkey Property in 1992, 1993, 1994, and 1995 and observed the following activities during that period that may have resulted in contaminant releases:

- Improper storage of sandblast grit, solvents, and paint sludge at a metalfabricating shop; and
- Debris and drums containing oily substances scattered around the industrial park.

#### 2.3.1.3 Operations on the Sesko Property

The Sesko Property was used for bulk petroleum storage and distribution from as early as 1946 to no later than 1993, when the aboveground storage tanks (ASTs) were removed. Lent's was the primary operator of the tank farm on the Sesko Property. Former AST locations are shown on Figure 2-4. A process flow diagram of petroleum storage and distribution operations is provided on Figure 2-5. Since 1993, the Sesko Property has been used for boat maintenance, automobile salvage, equipment and debris storage, parking, and metal reclamation. The owner of the Sesko Property was involved in legal disputes with the City over nonconforming use of the Sesko Property (as a junkyard), violations of the Shoreline Management Act, and, in 2003, improper decommissioning of an underground storage tank (UST). Ecology spill records also indicate that approximately 25 gallons of gasoline were released from the Sesko Property to surface water in January 2003. The majority of the equipment and debris has been removed, and the Sesko Property is currently vacant.

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<sup>&</sup>lt;sup>6</sup> Based on City directory information, Lent's continued operating on the McConkey Property for at least 3 years after the McConkeys acquired the majority of the McConkey Property in 1979.

The Sesko Property includes remnants of the Former Ravine, which has been filled over the years. Fill activities have included the following:

- Before 1930. No records documenting fill activities before operation of the
  former gas works have been identified. However, based on a comparison of the
  1919 shoreline (Figure 2-4) with an aerial photograph dated 1946 and sewer
  maps dated 1939, it appears that a portion of the Former Ravine was likely filled
  by the late 1930s, before construction of a historical residence located on the
  Sesko Property and before construction of the Lent's tank farm.
- 1931 to 1955. Aerial photographs and recorded observations (Tymstra 1942 and Judd 2014) indicate that the western portion of the Former Ravine was filled between 1931 and 1955. Recorded observations indicate that people unaffiliated with the former gas works dumped miscellaneous garbage, trash, and fill in the Former Ravine before 1942. Residual materials from former gas works operations (i.e., soot, ashes, cinders, and tar-laden wood chips and shavings) were also reportedly dumped in the Former Ravine during this period (see Section 2.3.1).
- 1941 to 1974. An easement granted by Western to the City gave the City the right to dump refuse, garbage, and ashes from an incinerator into the Former Ravine. The easement reserved the right for Western to dump ashes and cinders in the easement area, which included the eastern 25 feet of the Former Gas Works Property (most of which lies on the current Sesko Property). According to the City, the historical records that partially document this time period were destroyed in a fire, and any documents regarding construction of the incinerator or dumping of refuse, garbage, or incinerator ash into the Former Ravine would have been lost in that fire.
- **1968.** A DNR inspection reported that concrete and piping debris were placed in the Former Ravine (DNR 1968).

Petroleum transfer lines that connected a dock located on the north edge of the Sesko Property (Former Sesko Dock) to the Former ARCO Property and the Lent's tank farm were formerly located on the Sesko Property and may still be in place. An employee of the owner of the Sesko Property indicated that he had removed a portion of underground petroleum transfer piping he encountered in the northern portion of the Sesko Property. Petroleum transfer lines also reportedly connected the Former Sesko Dock to the Former SC Fuels Property to the east. Approximate pipeline locations, shown on Figure 2-4, were identified on construction plans for City sewer improvements (CH2MHill 1982; MH&A 1982).

#### 2.3.1.4 Historical Operations Data Needs and Collection Strategy

Uncertainties regarding historical operations of the former gas works include the following:

- Historical locations where tarry residuals were managed or placed; locations identified in historical records are approximate, and the specific locations of tar pits identified by Tymstra (1942) were not identified, although they may correspond with the tar pit identified by a former resident in the same time period (Judd 2014);
- Location of transfer lines from storage tanks to the generators; methods of conveyance/locations of pipelines of tar, oil, and gas liquor to residual management areas or byproduct storage tanks; and location of transfer lines from byproduct storage tanks to the Former Gas Works Dock; and
- Presence of subsurface structures (sumps, tar wells, and gas holder foundation) that may harbor process residuals.

Resolution of these uncertainties would assist in identifying locations of potential contaminant releases. Investigation methods to identify underground structures or former tar pits include geophysical survey and subsurface explorations such as test pits or trenches.

### 2.3.2 Adjoining Properties

Surrounding properties include: (1) the Penn Plaza Property, which is located to the south of the McConkey Property, (2) the Former ARCO Property, which is located to the west of the McConkey Property across Thompson Drive, and (3) the Former SC Fuels Property, which is located to the east of the Sesko Property across Pennsylvania Avenue (Figure 2-1). Historical and current operations on these properties are discussed in the following subsections.

#### 2.3.2.1 Penn Plaza Property

There are five buildings on the Penn Plaza Property, which is used as an industrial park. Multiple tenants occupy the industrial park. Based on available records, the Penn Plaza Property has been used for commercial and/or industrial uses since the late 1930s or early 1940s. Prior to this time, an intermittent stream ran northeast across the Penn Plaza Property toward the Former Ravine on the Sesko Property. This stream was reportedly used by area residents for dumping refuse and was filled in by 1942 (Judd 2014).

Operations on the Penn Plaza Property have included Lent's operations from the 1940s to approximately 1985 and industrial park operations from approximately 1985 to the

present. <sup>7</sup> Lent's operations on the Penn Plaza Property included spray painting, metal plating, a pipe shop, truck repair, and parking for petroleum distribution. <sup>8</sup> A former employee of Cascade, who worked in Bremerton in 1968 and 1969, recalled that wood treating may also have occurred as part of Lent's operations (Clapp 1997). Since the cessation of Lent's operations, multiple tenants have used the Penn Plaza Property for industrial uses, including sheet metal fabrication, floating pier and acrylic septic tank manufacturing, concrete pipe/manhole manufacturing, heating and air conditioning repair, and marine propeller repair (TechLaw 2006; Hart Crowser 2007).

Ecology inspected operations at the Penn Plaza Property in 1992, 1993, 1994, and 1995 and identified the following activities that may have resulted in contaminant releases:

- A tenant reported to Ecology that an electroplating operation had made illegal discharges to a storm drain that resulted in a sewer backup.
- Ecology observed improper storage of waste concrete and waste oil at one of the tenant locations.
- Ecology observed diesel staining on the ground at another tenant location.
- Ecology observed debris and drums containing oily substances scattered around the industrial park.

On the north end of the Penn Plaza Property are oil and gasoline supply pipelines that connected the Former Sesko Dock with the Former ARCO Property to the west. The approximate location of these pipelines, based on a utility locate conducted during the 2010 TCRA, is shown on Figure 2-4.

#### 2.3.2.2 Former ARCO Property

The Former ARCO Property was used for bulk petroleum storage and distribution from the mid-1940s to the late 1980s or early 1990s. Initially, 4 ASTs were present, with 2 added prior to 1956, 5 added in the late 1970s, and 4 added in the early 1980s for a total of 15 ASTs. Loading racks were located in the southeast corner of the Former ARCO Property. All tanks were removed by 1993. Property records indicate storage of gasoline, diesel, and oil. Product lines connected the ASTs on the Former ARCO Property with the Former Sesko Dock. Piping from the Former ARCO Property crossed the adjacent property to the north and ran west along the waterfront to a former dock (Former ARCO Dock) located approximately where the Port Washington Marina is today (see Section

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<sup>&</sup>lt;sup>7</sup> Based on City directory information, Lent's continued operating on the McConkey Property for at least 3 years after the property was sold in 1979.

<sup>8</sup> Petroleum for Lent's petroleum distribution was stored on what is now the Sesko Property.

2.3.3). According to a former resident, the piping to the Former ARCO Dock was located above ground (Judd 2014).

Since the early 1990s, the Former ARCO Property has been sporadically occupied by various tenants, including a tenant that conducted furniture refinishing and repair. The Former ARCO Property is currently being used for commercial use by Pipeworks Mechanical and Service, Inc.

#### 2.3.2.3 Former SC Fuels Property

The Former SC Fuels Property was used for bulk petroleum storage and distribution from the mid-1940s to the present. Operations on the Former SC Fuels Property are currently inactive. Initially, five ASTs were present, with one AST added prior to 1963, for a total of six ASTs. Four USTs were removed in 2003. Property records indicate storage of gasoline, diesel, and waste oil.

The Former SC Fuels Property is registered in Ecology's Voluntary Cleanup Program. A series of environmental investigations and remedial actions performed between 1997 and 2007 have confirmed releases of petroleum products and associated constituents, including gasoline, diesel, oil, BTEX, and PAHs. Additional information about the investigations and remedial actions is provided in Section 4.3.1.

Stormwater at the Former SC Fuels Property is collected in a series of catch basins, piped to an oil-water separator located at the top of the bluff, and discharged through an outfall to the Port Washington Narrows (Figure 2-4). Ecology conducted a site visit in 2006 and noted a "gasoline odor" along the shoreline of the Former SC Fuels Property close to the stormwater outfall.

Pipes supplying petroleum to the Former SC Fuels Property tank farm ran from the Former SC Fuels Dock (see Section 2.3.3). An unknown number of petroleum transfer pipes also reportedly ran from the Former Sesko Dock to the tank farm on the Former SC Fuels Property, although their alignment is unknown (see Section 2.3.1.3).

#### 2.3.2.4 Adjoining Properties Data Needs and Collection Strategy

The data needs associated with the adjoining properties consist of the following:

- Investigation of the drip tank associated with the former gas works distribution piping, which was located on the north end of the Penn Plaza Property, as a potential source of contamination; and
- Determining whether releases of hazardous substances that may have occurred on the adjoining properties may be migrating onto the Former Gas Works Property and commingling with gas works-related contamination.

### 2.3.3 Aquatic Parcels

Four docks were constructed in the aquatic parcels located adjacent (or closest to) to the properties described in Sections 2.3.1 and 2.3.2 (Figure 4-2). These aquatic parcels were

leased from DNR. A description and brief history of each dock is included in the following paragraphs, and a detailed lease history prepared by DNR is provided in Appendix B.

#### 2.3.3.1 Former Gas Works Dock

The Former Gas Works Dock was constructed by Western on November 25, 1930, as part of the development of the former gas works. It was located on the aquatic parcel adjacent and to the north of the Former Gas Works Property. The Former Gas Works Dock was used to offload coal, briquettes, and oil (via a 3-inch-diameter pipeline). Records indicate that the Former Gas Works Dock was also used to transfer heavy-end byproducts. In 1948, as part of the propane blending retrofit, the Former Gas Works Dock was updated to allow offloading of propane gas. Based on review of aerial photography, the Former Gas Works Dock was removed sometime between 1971 and 1974.

#### 2.3.3.2 Former ARCO Dock

The Former ARCO Dock was constructed by the Richfield Oil Corporation in approximately 1942. It was located on the aquatic parcel immediately adjacent and to the west of the aquatic parcel operated by the former gas works. The Former ARCO Dock served as both boat moorage and support for the pipelines associated with upland ARCO operations. It was removed by Richfield Oil's successor in the mid-1980s.

#### 2.3.3.3 Former Sesko Dock

The Former Sesko Dock was constructed by Lent's in approximately 1942. It was located on the aquatic parcel immediately adjacent and to the east of the aquatic parcel operated by the former gas works. The Former Sesko Dock was used to support supply pipelines for barge delivery of diesel and stove oil, which were stored on the Sesko Property. During the 1970s and 1980s, the Former Sesko Dock was also used to supply the tank farm on the Former ARCO Property and the tank farm on the Former SC Fuels Property. In 1993, the pipelines on the Former Sesko Dock were removed. The Former Sesko Dock was removed in September 2001 pursuant to a DNR order.

#### 2.3.3.4 Former SC Fuels Dock

The Former SC Fuels Dock was constructed by General Petroleum Corporation of California in 1942. It was located on the aquatic parcel immediately adjacent and to the east of the aquatic parcel where the Former Sesko Dock was located. The Former SC Fuels Dock was constructed for the purpose of handling petroleum products. The Former SC Fuels Dock was removed in 1967 by Mobil Oil Corporation when barge deliveries of petroleum products were discontinued.

### 3 Environmental Setting

### 3.1 Climate and Meteorology

The Bremerton, Washington, area is dominated by a marine temperate climate with cool and comparatively dry summers and mild, wet, and cloudy winters (WRCC 2014). The average annual high temperature for Bremerton is 60 degrees Fahrenheit (° F), and the average annual low temperature is 43° F (WRCC 2014). Average annual precipitation is 52 inches, with nearly half of that occurring in November, December, and January (WRCC 2014). During this wet season, rainfall is usually light to moderate in intensity and continuous over a period of time, rather than brief, heavy downpours. During the driest months of July and August, it is not unusual for 2 to 4 weeks to pass with only a few showers (WRCC 2014). The prevailing wind direction in the region is south or southwest during the wet season and northwest in summer, with an average wind velocity of less than 10 miles per hour (WRCC 2014).

### 3.2 Topography and Drainage

The Former Gas Works Property is located on a bluff on the south shore of the Port Washington Narrows. The Former Gas Works Property generally slopes gently to the north and is covered with buildings or pavement. At the northern edge of the Former Gas Works Property, a vegetated bluff slopes steeply down to the beach. Over time, the bluff has expanded to the north with the placement of fill material. Remains of the Former Ravine along the eastern edge of the Former Gas Works Property can be seen as a cove located at the northern edge of the Sesko Property. Stormwater drainage characteristics on the Former Gas Works Property and adjacent properties are as follows:

- McConkey and Penn Plaza Properties. Pavement covers most of the McConkey and Penn Plaza Properties, and the properties have catch basins connected to the City stormwater drainage system. A City stormwater and combined sewer overflow (CSO) outfall is located offshore, north of Pennsylvania Avenue. A catch basin in the northwest corner of the McConkey Property is connected to an outfall on the beach below the bluff.
- **Sesko Property.** Most of the Sesko Property is unpaved. Stormwater either infiltrates or runs off, presumably to the north toward the Port Washington Narrows.

### 3.3 Geology and Hydrogeology

### 3.3.1 Regional Geologic Setting

The Site lies within the Puget Lowland, an area that has alternated between glacial and interglacial environments during the last 2 million years. The result has been a stacked and imperfectly preserved sequence of glacial and nonglacial strata. This irregular

stratification has been further impacted by the tectonics of the Seattle fault, a regional thrust fault system that extends through the area, including a strand through Oyster Bay. The impacts of the fault system include uplift and tilting of bedrock and Quaternary strata in some areas and subsidence in others.

Interglacial climates produced sediments much like the forested Puget Lowland before extensive development, with broad floodplains and gently sloping uplands. These deposits include silty to sandy floodplain sediments, scattered gravelly channel deposits, and peat and lacustrine (lake) sediments. Glacial climates resulted in rapid accumulation of glacial sediments and scour of preexisting landforms and deposits. These deposits include advance glacial lake (glaciolacustrine) deposits, advance outwash (glacial river deposits), glacial till (subglacial deposits), and recessional glacial deposits.

Bedrock crops out on the northern end of the peninsulas between Phinney Bay and Ostrich Bay, and elsewhere generally north and west of the Site. Map data and limited deep well data suggest that bedrock generally dips to the south and west below the Site area. This bedrock dip forms a regional basement aquitard. Some of the older sediments above bedrock are also likely tipped in this direction due to regional rotation along the Seattle fault. Younger deposits, including those encountered in explorations for this project, are expected to be generally more horizontal but will include a number of discontinuous and irregularly shaped lenses of fine- and coarse-grained sediments that will impact the velocity and direction of groundwater flow. A conceptual geologic model of the Site area, including surficial geology (Figure 3-1) and subsurface geology (Cross Section AA–AA' on Figure 3-2) has been developed using regional map and well log data. Areas below the known exploration depths are shown as "undifferentiated."

The conceptual regional hydrogeologic model is one of rainfall and infiltration on an upland covered generally with till and glacial outwash. Some of this water runs off as stormwater, while a portion infiltrates. The water that infiltrates (groundwater) will migrate more quickly through more-permeable strata and will be generally retarded by less-permeable strata. The migration of water through these strata is influenced by the location and dip of the low-permeability strata (aquitards), as well as the location of waterways and other low-lying areas, which are often points of groundwater discharge. Regional patterns indicate that uplands are generally recharge areas, and slopes near sea level are discharge points. Groundwater also migrates from deeper strata and discharges upward into waterways.

### 3.3.2 Site Geology

Four principal geologic units have been identified based on previous explorations: fill, natural glacial deposits of the Vashon Drift, nonglacial deposits from one or more of the interglacial events that preceded the Vashon glaciation, and deposits from an older glaciation. The characteristics and distribution of these major sequences are described in this section, from the stratigraphic top (generally younger) to the bottom. Note that these geologic interpretations are based on logs prepared by multiple geologists over the

course of the prior investigations. Subsurface interpretations from these earlier explorations (e.g., fill characteristics or extent) may be refined later based on future observations.

The locations of the cross sections are shown on Figure 3-3, and four geologic cross sections, are provided on Figures 3-4 through 3-7. Soil boring logs are provided in Appendix C. A description of the soils observed at the Site is provided in the following text.

Although fill was not specifically identified in many of the soil boring logs, it was apparently present in the majority of the previous explorations at the Site, in thicknesses ranging from a foot or less to about 15 feet. The thickest fill is present in the Former Ravine area on the Sesko Property. Fill is generally composed of brown to black, loose to very dense, or stiff to very stiff variable mixtures of silt and sand with variable amounts of gravel, coal fragments, asphaltic concrete, and other debris. The density and consistency of the fill was generally high for nonstructurally placed fills and may be due to inclusion of ash in the fill soils, which can produce slight cementation of soils.

Over the majority of the Site, glacial deposits were encountered beneath the surficial fill. The geologic maps of the Site indicate the glacial unit is the Vashon Drift. The soils encountered in the explorations generally consisted of clean (fines are absent) to silty fine- to medium-grained sand with trace to minor amounts of gravel and scattered interbeds of sandy silt. These glacial deposits were observed to be dense to very dense and were generally brown to gray. The gradation and density of this unit suggests that it is primarily Vashon advance glacial outwash. This unit has moderate permeability and, where saturated, will form an aquifer.

Pre-Fraser nonglacial deposits (predating the Vashon Glaciation) are present in the bluffs and uplands in the northeastern portion of the Site. Explorations encountered olive to gray and brown, stiff to hard silt to sandy silt with interbeds of very dense silty sand. Thin interbeds or lenses of clay and silty clay and scattered gravelly layers may be present. This unit generally has low permeability; however, cleaner sandy layers may become saturated.

An older glacial sequence is present below the Vashon outwash and the pre-Fraser nonglacial deposits. The older glacial sequence consists of lenses or discontinuous layers of glacial till within an outwash-like brown to gray, very dense slightly silty to silty sand. The lenses of till are composed of brown to gray very dense silty gravel with sand and silty sand with gravel. The till lenses are generally considered an aquitard, but the outwash-like silty sand component was noted to be wet below about the 5 to 10 foot elevation, which probably reflects the regional water table. Additional investigations will be conducted to determine whether till acts as an aquitard at the Site, as described in Section 3.3.4.

### 3.3.3 Hydrogeology

Groundwater on the McConkey Property and Sesko Property was encountered at depths between 15 and 41 feet. Groundwater elevations have ranged between 3 and 10 feet above mean sea level, with an estimated flow direction to the north-northwest (to the Port Washington Narrows) during one sampling event (GeoEngineers 2007b). Monitoring well construction details and groundwater elevation measurements are summarized in Table 3-1. Well construction logs are included in Appendix C.

Groundwater on the Former SC Fuels Property has been encountered at depths between 4 and 15 feet, with an estimated flow direction to the northwest. Groundwater on the Former SC Fuels Property appears to be perched within sandy zones present in generally low-permeability nonglacial soils.

The estimated directions of groundwater flow on the McConkey, Sesko, and Former SC Fuels Properties, based on previous studies, are shown on Figure 3-8. However, groundwater studies to date have not evaluated the effect of tidal influence on-site groundwater levels and flow direction. One-time groundwater elevation measurements are prone to error if tidal effects are significant.

# 3.3.4 Geology and Hydrogeology Data Needs and Collection Strategy

Data needed to further characterize Site geology and hydrogeology include the following:

- Identification of aquifer zones impacted by Site contamination. This would be determined by soil and groundwater sampling to characterize subsurface lithology and determine the nature and extent of contamination (see Section 7).
- Identification of aquitards underlying or between impacted aquifer zones. This
  would be determined through a combination of data collection methods,
  including characterization of subsurface lithology, evaluation of physical soil
  characteristics, and evaluation of hydraulic conductivity and gradients.
- Soil characteristics of aquifer and aquitard materials, including grain size, density, porosity, and organic carbon content. A subset of soil samples collected for chemical analysis would be tested for physical parameters.
- Hydraulic conductivity of aquifer materials. This would be measured using slug testing of Site wells.
- Hydraulic gradients and groundwater flow direction, including characterization of tidal influences and seasonal variability. These would be measured using a network of pressure transducers installed in wells over tidal cycles and during different seasons.

### 3.4 Human Populations and Land Use

The Former Gas Works Property is located in Bremerton, which is the largest city on the Kitsap Peninsula and home to Puget Sound Naval Shipyard and the Bremerton Annex of Naval Kitsap Base. According to the 2010 census, the population of Bremerton is 37,729 people with 1,328 inhabitants per square mile. The racial makeup of Bremerton is predominantly white/Caucasian (74%) with the rest of the population classified as "other" or two or more races (10.4%), African American (6.7%), Asian (5.5%), Native American (2.0%), and Pacific Islander (1.3%)<sup>9</sup>. According to the 2000 census, the total population of the Suquamish Tribe is 616 people.

The Former Gas Works Property is in an area of industrial-zoned properties that includes the Former ARCO Property and Former SC Fuels Property. Surrounding this industrial property core are residential properties and a marina. A zoning map is included on Figure 2-1.

#### 3.4.1 Tribal Use

Tribal commercial, subsistence, and ceremonial fisheries have historically occurred in Dyes Inlet and the Port Washington Narrows. The Tribe has stated that "Suquamish tribal members fully intend to continue to fish these areas for cultural, subsistence and commercial purposes" (Suquamish Tribe 2014). "The Tribe uses the Washington Commercial Shellfish Growing Area Classification to determine the suitability of bivalve harvests (i.e., clams, oysters)" (Suquamish 2011). The marine area adjacent to the Former Gasworks Property is designated as "Unclassified," due to proximity to CSOs, which precludes shellfish harvesting. However, the harvest of finfish and other marine invertebrates (i.e., crab and sea cucumber) are not restricted adjacent to the Former Gas Works Property (Suquamish 2011).

### 3.4.2 Drinking Water Use

Water services at the Site and surrounding area are supplied by the City. The closest public water supply wells are located over one mile from the Site. The use of private wells within the Bremerton Water Service Area is not allowed, and there are no drinking water wells near the Site listed in Ecology's database.

<sup>&</sup>lt;sup>9</sup>All work conducted during the RI/FS will be conducted in a manner consistent with EPA's Environmental Justice principles. EPA defines Environmental Justice as: "The fair treatment and meaningful involvement of all peopleregardless of race, color, sex, national origin, or income with respect to the development, implementation and enforcement of environmental laws, regulations, and policies."

The Site is located adjacent to the Port Washington Narrows, a saltwater body. The extent of saltwater intrusion and the potability of Site groundwater and its potential future use as a drinking water source have not been evaluated.

### 3.4.3 Land Use Data Needs and Collection Strategy

Data needed to further characterize land use include the following:

 Evaluation of potential beneficial use of impacted aquifers. This would be evaluated by characterizing the extent of impacted groundwater and the physical and geochemical characteristics of the impacted aquifers, including hydraulic conductivity and salinity.

No data needs are currently identified for the use of the Port Washington Narrows. Through the RI/FS process, the Suquamish Tribe may provide additional information pertaining to historical and current tribal land use in the vicinity of the Site.

### 3.5 Port Washington Narrows and Dyes Inlet

The Former Gas Works Property is located along the Port Washington Narrows, which is a tidal channel connecting Dyes Inlet to Sinclair Inlet and Puget Sound. Dyes Inlet is a terminal estuary, comprising five embayments (Phinney, Mud, Ostrich, Oyster, and Chico Bays) and the Port Washington Narrows (Figure 3-9).

The waters of Port Washington Narrows are relatively shallow, with average depths of less than 30 feet. Depths within Dyes Inlet range up to 100 feet but are typically less than 50 feet. Area bathymetry is shown on Figure 3-9.

The shorelines of the Port Washington Narrows and Dyes Inlet have been extensively developed. These shorelines include the cities of Bremerton and Silverdale as well as the community of Tracyton. Other significant features include several former U.S. Navy facilities and regional transportation networks, including State Routes 3 and 303. The Warren Avenue and Manette Bridges are located across the Port Washington Narrows east of the Former Gas Works Property.

Hydrologic inputs to the Port Washington Narrows and Dyes Inlet include the tidal exchange with Sinclair Inlet and freshwater inflows from both stream and piped flows. Figure 3-9 summarizes compiled information from Kitsap County and the City regarding identified stormwater outfalls, CSO discharge points, and surface water inputs. Additional private and municipal outfalls may be present in addition to those identified by these information sources.

Hydraulic exchange between Dyes Inlet, the Port Washington Narrows, and the balance of Puget Sound is limited by the geography and the resulting hydrodynamics. In addition to tide and current data available from public sources (e.g., National Oceanographic and Atmospheric Administration [NOAA]), the waters of Dyes Inlet and the Port Washington Narrows have been studied as part of regional water quality programs. Total maximum

daily load studies and a contaminant mass balance evaluation have been performed for Dyes Inlet and may provide useful data for the RI/FS. Hydrodynamic modeling of the area has been performed as part of regional studies of Puget Sound. The results of additional studies are available to characterize environmental quality within Sinclair Inlet, immediately south of Dyes Inlet and the Port Washington Narrows. The Sinclair Inlet studies include extensive testing that has been performed in association with the Bremerton Naval Shipyard, as well as other regional study programs.

Additional data may be needed to evaluate the feasibility of remedial alternatives as discussed in Section 9.

#### 3.6 Natural Resources

This section describes the natural resources of the upland areas, aquatic habitats, and related data needs for the RI/FS.

### 3.6.1 Upland Areas

The upland areas of the Former Gas Works Property and surrounding areas have been developed for industrial uses consistent with zoning provisions. However, some terrestrial and riparian habitat is present, particularly on the bank adjacent to the Port Washington Narrows, the Former Ravine, and the shoreline areas of the McConkey and Sesko Properties. The Washington Department of Fish and Wildlife (WDFW) manages a Priority Habitats and Species Program (PHS). Preliminary queries of WDFW's PHS system did not identify any priority terrestrial natural resources on the parcels associated with the Former Gas Works Property.

### 3.6.2 Aquatic Habitats

Aquatic habitats at the Site include those in the beach and subtidal areas within and near the Former Gas Works Property. Shoreline and aquatic habitat adjacent to the Former Gas Works Property are located within the Tribe's Usual and Accustomed area. Fish and shellfish resources are present within the waters of the Port Washington Narrows and Dyes Inlet. Fish and crab are known to be present and support commercial, recreational, and tribal fisheries. Shellfish harvesting within the Port Washington Narrows and Dyes Inlet has been restricted due to water-quality-related shellfish harvesting closures. However, efforts have been made by state and local governments, tribes, and other stakeholders to improve water quality in the area and reduce or lift these shellfish harvesting restrictions. A number of shellfish enhancement projects have been proposed within portions of Dyes Inlet. It is not known what measures have been undertaken by the Washington State Department of Health or the Kitsap Public Health District (KPHD) to monitor illicit shellfish harvesting within Dyes Inlet or the intertidal areas adjacent to the Site. Signage indicating the closure of the beach adjacent to the Former Gas Works Property was installed as part of the 2013 TCRA (see Section 4.2.2).

The query of the WDFW PHS identified two aquatic natural resources in the vicinity of the Former Gas Works Property: estuarine intertidal aquatic habitat along the northern and southern shorelines of the Port Washington Narrows and hardshell clams along the northern shoreline of the Port Washington Narrows.

### 3.6.3 Natural Resources Data Needs and Collection Strategy

Natural resources at the Former Gas Works Property need to be further defined and delineated in order to plan and accurately conduct a risk assessment for the RI/FS. Terrestrial and aquatic natural resources at the Former Gas Works Property, including threatened and endangered species, will be documented and described as part of the RI. Natural resource information for terrestrial and aquatic habitats at the Site will be developed based on information in scientific literature, data compilations from state and federal agencies, and information from the Suquamish Tribe, as well as Site surveys to be conducted during the RI/FS. This work will include an evaluation of shellfish harvesting.

### 3.7 Cultural Resources

This section describes the archaeological sites and historic structures at and around the Site and discusses the cultural resources data needs and collection strategy.

### 3.7.1 Archaeological Sites and Historic Structures

There are no recorded archaeological sites or historic structures at the Former Gas Works Property or in the immediate vicinity. However, no cultural resources surveys have been conducted on the Site or in the vicinity prior to the present project. The documented archaeological sites nearest to the Former Gas Works Property include the following:

- Site 45KP121, a pre-contact and historic-era shell midden site, is located in Evergreen Park, approximately 0.6 miles east-southeast of the project area;
- The Manette Site (45KP009), a large pre-contact midden and possible fortification site where human remains have reportedly been found, is located on a bluff above the beach, just west of the Manette Bridge (1.2 miles east-southeast of the project area); and
- A number of ethnographic place names have been recorded at various locations along the Port Washington Narrows.

Kitsap County assessor's records (accessed January 2014) indicate that there is one building older than 50 years on the Penn Plaza Property—a warehouse constructed in 1955. The structure has not been evaluated for National Register of Historic Places (NRHP) eligibility. No impacts on this structure are anticipated during the RI/FS.

An Anchor QEA archaeologist visited the project area in August 2013 to make a preliminary assessment of current conditions. The project area has been extensively modified in the historic and modern eras, with placement of fill materials and debris, and development and redevelopment of the Site for industrial uses. No native sediments, other than active beach deposits, were visible in the project area.

### 3.7.2 Cultural Resources Data Needs and Collection Strategy

Despite historical disturbance and filling activity at the Site, the presence of documented archaeological sites within the vicinity indicates that there remains some potential for archaeological resources to be present in native upland soils beneath the fill deposits. RI/FS activities that penetrate these native soils will incorporate appropriate measures to protect potential archaeological resources, including potential archaeological monitoring and implementation of an incidental discovery plan. RI/FS activities will include coordination with the Suquamish Tribe and/or the State of Washington, consistent with the requirements of Section 106 of the National Historic Preservation Act.

# 4 Previous Investigations and Remedial Actions

# 4.1 Initial Study Area Investigations

Previous environmental field investigations at the Former Gas Works Property include the following:

- Sesko Property Field Inspection (Ecology 1995);
- Preliminary Upland Assessment, McConkey and Sesko Properties (GeoEngineers 2007b); and
- Targeted Brownfields Assessment (TBA), McConkey and Sesko Properties (E&E 2009).

The upland exploration locations and sampling depths by analyte group are provided on Figure 4-1. The scope and general conclusions of each study are described in the following subsections.

### 4.1.1 Ecology Field Inspection (1995)

In 1995, Ecology collected three surface soil samples from the Sesko Property and one surface sediment sample from the tidelands just north of the Sesko Property. Samples were analyzed for metals and SVOCs. High concentrations of PAHs were detected. Ecology used the data in conducting a Site Hazard Assessment and ranked the Site a "1" (highest concern).

# 4.1.2 Preliminary Upland Assessment (2007)

In 2007, on behalf of the City and funded by a brownfield grant from EPA, GeoEngineers conducted a preliminary assessment of the McConkey and Sesko Properties (GeoEngineers 2007a) that included the following:

- Advancing eight soil borings and collecting soil samples to a maximum depth of 45 feet;
- Installing monitoring wells at each of the eight soil boring locations and collecting groundwater samples; and
- Analyzing soil and groundwater samples for total petroleum hydrocarbons (TPH),
   VOCs, SVOCs, polychlorinated biphenyls (PCBs), and metals.

This work identified relatively high concentrations of gasoline- and diesel-range TPH, VOCs including benzene, and PAHs in soil and groundwater on the McConkey and Sesko Properties. VOCs and PAHs were detected in soil samples at depths up to 35 feet. Several metals, including arsenic, lead, and chromium (including chromium VI), were detected in groundwater at concentrations above potential drinking water cleanup standards.

### 4.1.3 Targeted Brownfield Assessment (2008)

In 2008, on behalf of EPA, E&E conducted a TBA of the McConkey and Sesko Properties (E&E 2008) that included the following:

- Advancing seven soil borings and collecting soil samples to a maximum depth of 45 feet;
- Installing monitoring wells at two of the seven boring locations;
- Collecting groundwater samples from the two wells and from temporary screens placed at four of the seven soil boring locations;
- Collecting five surface sediment samples from the beach north of the properties;
- Analyzing soil, groundwater, and sediment samples for TPH, VOCs, SVOCs, and metals.

Similar to the Preliminary Upland Assessment, this work identified relatively high concentrations of gasoline- and diesel-range TPH, VOCs including benzene, and PAHs in soil and groundwater on the McConkey and Sesko Properties. The assessment also identified relatively high concentrations of PAHs in surface sediments. VOCs and PAHs were detected in soil samples at depths up to 45 feet.

# 4.2 Initial Study Area Removal Actions

### 4.2.1 Time Critical Removal Action (2010)

In August 2010, sheens on the surface water of the Port Washington Narrows were reported to KPHD. Upon further investigation, KPHD identified a 12-inch-diameter concrete pipe that appeared to be the source of the sheen. The pipe is believed to be an abandoned City CSO outfall. KPHD reported the release to EPA, which in turn notified USCG for a response because the pipe was within its jurisdiction. In 2010, at the request of EPA, Ecology & Environment, Inc. (E&E) conducted sampling and analysis as part of the EPA and USCG's initial response. The response sampling included the collection of 32 surface sediment samples from a depth of 0 to 6 inches. The sediment samples were analyzed for VOCs and SVOC, both of which were detected in some samples.

EPA, DNR, KPHD, and Ecology entered into a USCG-led coordinated response under a Unified Command Structure. Cascade became aware of the response in October of 2010 and informed the USCG that it was interested in contributing to the response. USCG subsequently added Cascade to the Unified Command Structure and issued Cascade an Administrative Order for a Pollution Incident (Order) to implement response actions at the Site under the oversight of USCG. Cascade accepted the Order in a letter dated October 29, 2010.

In response to the Order, Cascade developed a Work Plan for the Incident Action and 2010 TCRA (Anchor QEA and Aspect Consulting 2010), which outlined the scope and details of the 2010 TCRA. The 2010 TCRA included the following key elements:

- Investigation of the location and orientation of the abandoned pipe;
- Permanent plugging of the pipe as close as practicable to the shoreline;
- Removal of all portions of the pipe from the new plug to the terminus of the pipe;
- Backfilling of the excavation created by removal of the pipe with clean beach material;
- Placement of an organoclay mat over impacted sediments (with minimal disturbance) near the terminus of the pipe that were observed to generate sheen; and
- Continued maintenance of a containment system until field observations and inspections confirm the situation is stable (no sheen).

On November 5, 2010, USCG and the other members of the Unified Command Structure approved the Work Plan. Cascade commenced the TCRA immediately upon approval and completed the 2010 TCRA on November 8, 2010 (Anchor QEA 2011). The Removal Action satisfied the following objectives of the Work Plan:

- The pipe was located and traced to the shoreline.
- The pipe was plugged as close as practicable to the shoreline, at the location specified in the Work Plan.
- All pipe sections downgradient of the new plug were removed together with all overburden sediments.
- All excavations were filled to grade with clean beach material.
- The organoclay mat was placed over the area of impacted sediments specified in the Work Plan.

Inspections of the 2010 TCRA area were completed as specified in the Work Plan. No surficial sheens related to the 2010 TCRA have been observed to date. Figure 4-2 shows the constructed elements of the 2010 TCRA.

# 4.2.2 Time Critical Removal Action (2013)

In 2013, Cascade completed a Removal Evaluation pursuant to the requirements of the AOC and the EPA-approved Removal Evaluation Work Plan (Anchor QEA and Aspect Consulting 2013a). The objective of the Removal Evaluation was to assess whether suspected migration pathways at the Site pose a threat to human health, welfare, or the environment if left unaddressed before completion of the RI/FS. The results of the

Removal Evaluation were reported in the EPA-approved Removal Evaluation Report (Anchor QEA and Aspect Consulting 2013c). The Removal Evaluation identified the following conditions that warranted action before completion of the RI/FS:

- Stormwater intrusion into Manhole A. Manhole A was believed to remain connected to the 12-inch-diameter concrete pipe that was plugged as part of the 2010 TCRA. Based on inspections conducted as part of the Removal Evaluation, stormwater could have been entering Manhole A through surface runoff or via a piping connection to Manhole A from a nearby sump. Stormwater entering Manhole A posed a risk of hydraulically surcharging the pipe plugged during the 2010 TCRA, which in turn could have increased the risk of a hazardous substances release to the Port Washington Narrows.
- Hydrocarbon sheen and deposits of solid hydrocarbon material in the SG-04/SG-05 area. Hydrocarbon sheens were observed in shallow subsurface sediments in the western area of the beach, near sampling stations SG-04 and SG-05. Surficial solid hydrocarbon material was also observed in the SG-04/SG-05 area. Both the sediments containing hydrocarbon sheen and the solid hydrocarbon material contained concentrations of PAH compounds that were elevated in comparison to those of the surrounding beach sediments.

The Removal Evaluation Report proposed the following removal actions in response to the identified conditions:

- Plug the connections to Manhole A. This action was intended to minimize the risk of hydraulic surcharge to the pipe plug and thereby minimize the risk of hydrocarbon releases from the pipe.
- Remove the accessible solid hydrocarbon material and place a cap over the sediments containing hydrocarbon sheen in the SG-04/SG-05 area. These actions were intended to minimize the risk of additional releases of hydrocarbons from this area to surface waters of the Port Washington Narrows and to prevent direct contact with these materials by beach users.
- Install signage. The purpose of the signs is to warn beach users regarding the presence of hydrocarbon contaminants in the beach sediments and provide agency contact information regarding the Site and the ongoing RI/FS process.

Upon completion of the Removal Evaluation, Cascade prepared a Removal Action Work Plan describing the proposed removal actions in more detail (Anchor QEA and Aspect Consulting 2013b). EPA approved the Final Work Plan and directed Cascade to perform the proposed removal actions (EPA 2013c). After EPA's approval, Cascade implemented the Removal Action (2013 TCRA), which met all of the objectives specified in the Work Plan including the following:

• Removing solid hydrocarbon material identified in the western beach area;

- Installing an organoclay mat and cover over the hydrocarbon sheen in subsurface sediments in the western beach area;
- Plugging Manhole A and the sump drain from the tank containment area;
- Completing beach monitoring inspections to confirm the effectiveness of the 2013 TCRA. Quarterly monitoring inspections are ongoing; and
- Installing required signage.

The work was completed in general accordance with the Work Plan and documented in the Removal Action Report (Anchor QEA and Aspect 2014). Three modifications to the scope of work specified in the Work Plan were made with EPA approval based on the observed conditions:

- The organoclay mat and cover in the northeastern portion of the designed mat and cover area was extended to cover sediments exposed by the removal of the solid hydrocarbon material from the intertidal area.
- Manhole A was plugged by means of a concrete ring extending above the ground surface capped with a bolted steel cover.
- Consistent with approvals from the City and pursuant to an access agreement with Penn Plaza Storage LLC, a catch basin draining into the tank containment area was rerouted to a City storm drain line to prevent accumulation of stormwater in the containment area.

Figure 4-2 shows the constructed elements of the 2013 TCRA.

### 4.3 Other Upland Investigations and Remedial Actions

This section describes work that has been conducted outside of the ISA (see Section 8.2) that is potentially relevant for characterizing Site and area-wide conditions.

# 4.3.1 Former SC Fuels Property Investigations and Remedial Actions (1997 to 2007)

Between 1997 and 2007, various consultants have performed soil and groundwater sampling at the Former SC Fuels Property (Pacific Environmental 1997; Noll 1999 and 2000; GeoEngineers 2002 and 2003; and GeoScience Management 2007), including the following:

- Advancing 13 hand-auger borings, 18 direct-push soil borings, and 15 hollowstem-auger borings to a maximum depth of 22 feet;
- Installing 15 monitoring wells to a maximum depth of 20 feet;
- Collecting 12 soil confirmation samples during removal of four USTs; and
- Analyzing soil and groundwater samples for TPH, BTEX, and/or lead.

The investigations indicated the presence of TPH and BTEX in soil and groundwater on the Former SC Fuels Property and in the eastern portion of the Pennsylvania Avenue right-of-way. The TPH and BTEX concentrations exceeded Washington State Model Toxics Control Act (MTCA) Method A cleanup levels.

# 4.4 Other Sediment Investigations and Remedial Actions

In addition to the sediment data developed as part of previous investigations and removal actions at the Site, other data sets have been compiled and studies completed within the Port Washington Narrows and Dyes Inlet that may provide information relevant to the RI/FS. Studies identified to date for these areas include the following:

- Chemical testing of sediments:
  - 2008 and 2009 Puget Sound Assessment and Monitoring Program
     (PSAMP 2005 and 2009) Spatial/Temporal Monitoring, Central Sound;
  - 1989 to 2013 PSAMP Long-Term/Temporal Monitoring (PSAMP 2005 and 2011a);
  - 2009 PSAMP Urban Waters Initiative, Bainbridge Basin (PSAMP 2005, 2009, and 2011b); and
  - 2009 Ocean Survey Vessel Bold Summer 2008 Survey Data Report (USACE 2009).
- Chemical testing of fish or shellfish tissue:
  - 2010 and 2012 Environmental Investment Project (ENVVEST) (Johnston et al. 2010; Brandenberger et al. 2012);
  - 2005 and 2007 NOAA Mussel Watch at station SIWP (NOAA 1993, 2006a, 2006b, and 2008); and
  - 2001 303d Ecology clam and crab sampling data (Ecology 2002).
- Studies of surface water quality:
  - An Integrated Watershed and Receiving Water Model for Fecal Coliform Fate and Transport in Sinclair and Dyes Inlets, Puget Sound, Washington (Johnston et al. 2009); and
  - Sinclair and Dyes Inlets Fecal Coliform Total Maximum Daily Load: TMDL and Water Quality Implementation Plan (Lawrence et al. 2012).
- Regional studies of contaminant source inputs to these water bodies:
  - Contaminant Mass Balance for Sinclair and Dyes Inlets, Puget Sound, Washington (Crecelius et al. 2003).

Additional studies may be identified later, during development of the RI/FS Work Plan and/or the implementation of the RI/FS. The evaluation of the above listed sediment and tissue data is discussed further in Section 7.

# 5 Preliminary Conceptual Site Model

This section presents a CSM based on available historical information, the current understanding of the environmental setting, and the findings of previous investigations (see Sections 2, 3, and 4). The CSM is a description of environmental conditions that includes sources of contamination, contaminant fate and transport in Site media, and potential routes of contaminant exposure for human and environmental receptors. A three-dimensional graphical CSM illustrating representative potential historical sources and migration of contaminants at the Site is shown on Figure 5-1. The nature and extent of specific contaminants is described in Section 7. The CSM will be further developed in the RI/FS Work Plan and during the RI and risk assessment as more Site-related information and data are gathered.

### 5.1 Potential Sources of Contamination

This section summarizes potential sources of contamination on the Former Gas Works Property and on surrounding properties. The potential sources and locations associated with known and documented operations (both MGP and other) are presented in the following sections; however, this discussion does not include undocumented or currently unknown potential source(s)/source areas, which may be identified through the collection and evaluation of data during the RI.

### 5.1.1 Former Gas Works Property Sources

Potential sources of contamination on the Former Gas Works Property include historical activities associated with the former gas works, as well as other activities on the Former Gas Works Property but unrelated to gas works operations.

### 5.1.1.1 Gas Works Operations

The potential primary sources associated with the production of manufactured gas are depicted on Figure 2-3. The area where the gas production process occurred is divided into potential source areas based on the predominant use and subsequent primary potential release mechanisms associated with each area. The primary potential source areas include the following:

- Coal/Coke Briquettes Area. As described in Section 2, solid feedstocks (coal and coke briquettes) were transported to the Former Gas Works Property by barge and offloaded and transported over the water, beach, and bluff to a concrete surface storage area in the northwest corner of the Former Gas Works Property. Coke briquettes have been observed on the beach and bluff, suggesting spills during the transport process. Additionally, coal/coke dust may have been swept off the concrete storage slab onto the surrounding ground surface.
- Tar and Petroleum Transfer Area. Petroleum products were delivered to the Former Gas Works Property and tar was removed from the Former Gas Works

Property by barge. Petroleum and tar from pipelines along the dock and at the connection to the barges may have been released directly to sediment or surface water. A pipeline presumably ran between the dock and the byproduct storage area to transport tar to the dock, but the location is unknown.

- Petroleum Storage Area. Petroleum products were stored in ASTs in the northeastern portion of the Former Gas Works Property. The products reported to have been stored in these tanks include gasoline and diesel fuel oil. Transfer piping presumably ran from the storage tanks to the furnaces, but the exact location of transfer piping is unknown. Petroleum may have been released from tanks and piping to soil at the surface or shallow subsurface in this area.
- Gas Generation and Purification Area. The main process area was located in the central portion of the Former Gas Works Property and included the furnaces, scrubber, gas holder, and purifier. The primary potential sources associated with the gas works process consist of spills, drips, and leaks of spent liquids, oils, gas liquor, tar, and tar-water mixtures from aboveground equipment, piping, and storage tanks to the ground surface.
- Residuals Management Area. A map of the former plant shows tar wells and a residue cistern to the east of the purifiers. These were likely used for separation of tar-water emulsions prior to resale of the tar. The details of the tar wells and residue cistern are unknown, but they likely extended into the shallow subsurface and may have been either lined or unlined at the base. A second area south of the main plant building was reportedly used for storage and/or separation of tar and tar-water emulsions in a tar pit. Oils and tar may have been released to the surface around these features or the subsurface beneath them.
- Tar and Light Oil Storage Area. The southern portion of the Former Gas Works Property was used for the storage of tar and light oil in ASTs. Tar and light oil may have leaked or been spilled onto the ground surface in the vicinity of the ASTs. Finished gas may have contained small amounts of oil that condensed in the distribution piping and were collected in the drip tank. Light oil may have been released to the shallow subsurface soil in the vicinity of the pipes and tank.
- Former Drainage Line Area. During the 2010 TCRA, a former drainage line on the Sesko Property that discharged to the Port Washington Narrows was identified. Tar-like hydrocarbons were identified in this drainage line, which was plugged during the 2010 TCRA (see Section 4.2.1). The drainage line is consistent with a former City CSO outfall documented in historical files. Wastewater and associated contaminants may have been discharged from this drainage line during operation of the former gas works.
- Ravine Fill Area and Shoreline Fill Area. Historical documents reference the surface disposal of gas works byproducts into the western portion of the Former

Ravine, to the east of the gas generation and purification area, and along the bluff to the north of the gas generation and purification area. Materials that were reportedly placed along the shoreline include ash, cinders, slag, and soot. Materials that were reportedly placed in the Former Ravine include ash, cinders, slag, soot, spent scrubber media (tar-laden wood chips and shavings), and spent purifier filter media (wood chips and/or iron oxide). Approximate areas of potentially gas-works-related fill are shown on Figure 2-3.

### 5.1.1.2 Other Operations

Other potential primary sources are associated with activities conducted after the shutdown and demolition of the former gas works, or they were conducted in the immediate vicinity of the former gas works. These sources are depicted on Figure 2-4 and summarized as follows:

- Bulk Petroleum Storage. Petroleum products were delivered to Lent's at a dock
  offshore of the Sesko Property and stored in ASTs for distribution by fuel delivery
  vehicles. Petroleum may have been released from piping and storage tanks to the
  ground surface and/or the shallow subsurface.
- Varied Light Industrial Use. Since the shutdown of the former gas works, the
  McConkey Property has been used for miscellaneous light industrial activities,
  including vehicle parking, metals fabrication, and equipment storage. Ecology site
  inspections in 1992, 1993, and 1994 indicated poor housekeeping practices
  associated with some of these operations. These operations are potential sources
  of solvents, metals, and petroleum hydrocarbons, which may have been released
  to the ground surface as either solids (sandblast grit, paint sludges, etc.) or
  components of liquids.
- Equipment Storage and Repair and Debris Filling. In addition to the bulk
  petroleum storage described above, activities on the Sesko Property since the
  shutdown of the former gas works include boat maintenance and storage,
  automobile salvage, and equipment and debris storage. These activities may be
  sources of contaminants to soil, sediment, and surface water by direct discharge,
  dumping, or spills to the ground surface.
- Other Operations. Other operations have reportedly included filling of the Former Ravine and shoreline areas, particularly on the Sesko Property. These operations may have included disposal of incinerator refuse, garbage, and ashes; placement of concrete and piping debris; and/or placement of miscellaneous metal, concrete, and fiberglass debris associated with maintenance and salvage of boats and equipment. Fill placed along the shoreline and in the Former Ravine may have included materials that contained hazardous substances. Although the presence of fill material alone does not necessarily represent a contaminant

source, hazardous substances associated with the fill may subsequently migrate to surrounding subsurface soil or groundwater.

### 5.1.1.3 Stormwater Discharge

Stormwater discharging to the Port Washington Narrows may contain contaminants and is a potential source of contamination to sediments or surface water. The outfalls that historically or currently capture water at the Former Gas Works Property are the following:

- Historical City Stormwater/CSO Outfall. As noted in Section 5.1.1.1, a historical drainage line and outfall were located within and offshore of the Sesko Property. A section of the drainage line on the beach was reportedly removed by the City during installation of a force main in the 1990s. The drainage line was plugged and partially removed as part of the 2010 TCRA (see Section 4.2.1). An upland manhole and storm drainage lines believed to be connected historically to the drainage line were plugged as part of the 2013 TCRA.
- McConkey Drainage Line. A small drainage line discharges stormwater from a shallow catch basin on the McConkey Property to the Port Washington Narrows.

### 5.1.2 Other Operations Sources – Adjacent Properties

Potential primary sources on adjacent properties include the following:

- Bulk Petroleum Storage. Petroleum products were delivered to bulk fuel storage
  facilities by barge at the Former ARCO Dock, the Former Sesko Dock, and the
  former SC Fuels Dock and stored in ASTs or USTs for distribution by fuel delivery
  vehicles. These petroleum storage facilities included the Former ARCO Property
  located west of the former gas works and the Former SC Fuels Property.
  Petroleum may have been released from piping and storage tanks to the ground
  surface and/or the shallow subsurface while these operations were ongoing.
- Varied Light Industrial Use. The Penn Plaza Property has been used for miscellaneous light industrial activities, including spray painting, a pipe shop, vehicle parking for a petroleum distributor, truck repair electroplating, metals fabrication, and equipment storage. Ecology site inspections in 1992, 1993, and 1994 indicated poor housekeeping practices associated with some of these activities. These activities are potential sources of solvents, metals, and petroleum hydrocarbons, which may have been released to the ground surface as either solids (sandblast grit, paint sludges, etc.) or components of liquids.

#### 5.1.2.1 Stormwater Discharge

As described in Section 3.5, a large number of documented stormwater and CSO outfalls are located within the Port Washington Narrows and Dyes Inlet, including the two

outfalls described in Section 5.1.1.3. Other nearby outfalls or discharge lines include the following:

- Current City Stormwater/CSO Outfall. An active City stormwater/CSO outfall is located along the Port Washington Narrows, offshore of the end of Pennsylvania Avenue. This outfall is located immediately adjacent to the 2010 TCRA area (Figure 4-2).
- Drain Line. A drain line from an oil-water separator on the Former SC Fuels Property discharges to the Port Washington Narrows.

# 5.2 Contaminant Migration and Transformation

Contaminants derived from the sources described in Section 5.1 may have been released to soil (surface and shallow subsurface), sediment, and/or surface water. Representative potential releases (e.g., leaks or spills from equipment, tanks, or piping; placement of contaminated fill materials; and discharges from outfalls) are shown conceptually on Figure 5-1. The released contaminants may have migrated from one location to another or from one medium to another. Contaminants may also undergo attenuation or transformation processes within media. The contaminant migration pathways and transformation processes are described in the following subsections.

### 5.2.1 Migration Pathways

Examples of potential contaminant migration pathways between media are shown conceptually on Figures 5.2, 5.3, and 5.4 and include the following:

- Migration of contaminants from surface soil to subsurface soil (e.g., leaching or product migration);
- Contaminant leaching or NAPL migration from soil/NAPL to groundwater;
- Groundwater/NAPL transport within the saturated zone;
- Groundwater discharges to surface water;
- Contaminant partitioning between groundwater and sediments (including sediment porewater);
- Migration of volatile NAPL/soil/groundwater contaminants to air;
- Migration of surface soil contaminants as fugitive dust;
- Release of surface soil contaminants to stormwater;
- Uptake of contaminants by terrestrial or aquatic biota; and
- Migration of contaminated sediments by sediment transport.

Based on the data collected to date (see Section 7), contaminants have been identified in soil, groundwater, and sediment. No Site-specific surface water, air, or tissue data are

available. Contaminant occurrences in these media may be due to direct releases or subsequent migration, for instance:

- Soil contamination may be the result of contaminated fill materials, downward flows of NAPL releases<sup>10</sup> through the subsurface and the coating of soil grains, or sorption of contaminants from other media (e.g., soil vapor, infiltrating stormwater, or groundwater).
- Groundwater contamination may be the result of direct discharge of contaminated aqueous materials and their migration downward through the subsurface and mixing with groundwater, leaching of NAPL in contact with groundwater, or stormwater infiltration of the subsurface, leaching of contaminants from NAPL or contaminated soil, and contaminant mixing with groundwater).
- Contaminants in sediment may be the result of direct releases to surface sediments (e.g., documented discharges from outfalls, undocumented spills, or leaks from dock piping and transfer operations); subsurface migration of contaminated groundwater or NAPL from the uplands, and migration through sediments; or a combination of sources. In particular, two sediment "hot-spot" areas were addressed by the 2010 and 2013 TCRAs:
  - The 2010 TCRA addressed a drainpipe that contained residual NAPL and surrounding contaminated sediments, which appeared to be the primary source of contamination in this area. The historical and ongoing contribution to sediment contamination from other potential sources in this area, including groundwater discharge, stormwater runoff, and the City CSO, is unknown.
  - The 2013 TCRA addressed an area of heavy sheen located in shallow subsurface sediments and solid surficial material containing high PAH concentrations. It is likely that the solid surficial material, which would be immobile in the subsurface, was placed at or near its locations; however, the source of the material is unknown. The source of the subsurface sheen is also unknown. During the TCRA investigation, a sheen was observed up to the base of the bluff. However, there are insufficient data to determine whether this contamination is contiguous with contamination in the upland.

<sup>&</sup>lt;sup>10</sup> Liquid releases generally will move downward, through the subsurface by means of gravity, but they may move laterally by preferential migration pathways if a barrier (e.g., low-permeability soils or, for NAPLs that are less dense than water, groundwater) is encountered.

Representative migration pathways, including subsurface migration pathways, are included on Figure 5-1.

### 5.2.2 Transformation Processes

In addition to contaminant migration pathways, contaminant concentrations in media can be reduced or attenuated by various combinations of natural processes. Examples of such processes include the following:

- Chemical or biological degradation of contaminants in soils, groundwater, sediments;
- Tidally induced mixing of groundwater near the groundwater/surface water interface;
- Natural recovery of marine sediments by burial, mixing, and/or degradation processes; and
- Metabolic transformation or elimination of chemical contaminants from the tissues of upland or aquatic biota.

# 5.2.3 Contaminant Transport and Transformation Data Needs and Collection Strategy

Additional data are needed to determine to what extent contaminants are migrating or could migrate in the subsurface. Sufficient data should be collected to aid in the assessment of contaminant transport. In particular, the data needs include characterization of the following:

- Soil lithology to identify potential subsurface migration pathways;
- Groundwater parameters governing transport rates and pathways (e.g., gradients and hydraulic conductivity to determine velocity; tidal effects; and salinity);
- Properties and extent of NAPLs in the subsurface;
- Extent of contaminants in environmental media;
- Groundwater chemistry along contaminant flow paths;
- Soil organic carbon content to evaluate leaching and sorption; and
- Natural attenuation parameters.

# 5.3 Exposure Pathways and Receptors

Exposure pathways and receptors that may be most relevant to the RI and risk assessment are summarized on Figures 5-3, 5-4, and 5-5. These figures illustrate how certain human and ecological receptors may use the Site and the impacted media that they could reasonably contact. The information included in this section will be further expanded during development of the RI/FS Work Plan, including a more exhaustive

review of human and fish/wildlife uses of the Site and vicinity and the rationale for focusing the risk assessment activities on the receptors and pathways summarized in Figures 5-3, 5-4, and 5-5.

Figure 5-3 illustrates different exposure pathways that could affect people using the Site or nearby areas. The potential exposure of people to Site-related contaminants of concern (COCs) differs in terms of both how those people use the Site and which areas of the Site are used. (i.e., beach/aquatic areas and upland areas). Some land uses could also change over time. For example, the Site is not zoned for residential land use, but as part of risk assessment activities, it may be prudent to evaluate potential future residential land use to understand the implications of changes in land use or zoning. Similarly, shellfish harvesting in the Port Washington Narrows is restricted due to shellfish harvesting closures unassociated with the former gas works. However, it may be prudent to evaluate potential future shellfish harvesting to understand potential exposures should those shellfish harvesting restrictions be lifted.

Preliminary complete current and future human exposure pathways to contaminated media include dermal contact with and incidental ingestion of soil, inhalation of fugitive dust and vapors, and consumption of fish/shellfish that are potentially contaminated with bioavailable Site-related contaminants. Preliminary incomplete current and future human exposure pathways will be further evaluated as part of the RI. The preliminary human exposure scenarios relevant to the Site include the following:

- Human Use of Beach/Aquatic Site Areas:
  - Recreational Beach Users. The potential for limited recreational beach use exists for individuals residing in proximity to the Site. During recreational use of the beach these individuals may be exposed to Site sediment and surface water.
  - Consumers of Fish/Crab from the Port Washington Narrows. The portions of the Port Washington Narrows adjacent to the Former Gas Works Property currently support the collection and consumption of fish and crabs under WDFW regulations. The Port Washington Narrows is also a Usual and Accustomed area of the Suquamish Tribe. Consumers of fish and crabs may also be exposed through incidental sediment and surface water ingestion during harvesting activities.
  - Consumers of Shellfish at the Site (Currently Restricted by Shellfish Harvesting Closures). The portions of the Port Washington Narrows adjacent to the Former Gas Works Property are currently closed to shellfish harvesting (due to water quality concerns associated with CSOs and other non-Site-related concerns) by Washington State Department of Health; however, exposures associated with shellfish harvesting will be evaluated to understand potential risks should the shellfish harvest

- restrictions be lifted. Consumers of shellfish may also be exposed through incidental sediment and surface water ingestion during harvesting activities.
- Beach Construction/Excavation Workers. This scenario relates to workers performing utility upgrades or maintenance or other activities that involve the disturbance of the beach/aquatic areas adjacent to the Former Gas Works Property. Beach construction workers could be exposed to Site surface and subsurface beach sediment.

### • Human Use of Upland Site Areas:

- Occupational Workers. The Former Gas Works Property and the properties in the vicinity are zoned for industrial uses. Occupational workers at the Site could be exposed to Site surface soil and vapor.
- Upland Construction/Excavation Workers. This scenario relates to
  workers performing utility upgrades or maintenance or other activities
  that involve the disturbance of soil at the Former Gas Works Property and
  the properties in the vicinity. Upland construction workers could be
  exposed to Site surface and subsurface soils and vapor.
- O Potential Future Residential Users of the Site (Not a Current or Planned Use). The Former Gas Works Property and the properties in the vicinity are zoned for industrial uses; and this is expected to remain the case for the foreseeable future. However, the potential for exposures of future residents may be appropriate to evaluate as part of the risk assessment to understand potential implications should property use be converted to residential. On-site residents could be exposed to Site surface soil and vapor. Although no water supply wells are located on or near the Former Gas Works Property, consumption of groundwater is retained as a potential pathway for screening, pending further evaluation of groundwater beneficial uses.

Preliminary complete aquatic-dependent ecological exposure pathways to contaminated media include direct contact with and ingestion of sediment, porewater, and marine water; and consumption of benthic invertebrates, fish, and other potentially contaminated prey. The risk assessment will include an evaluation of aquatic receptors with differing modes of exposure. Preliminary incomplete aquatic ecological receptors will be further evaluated as part of the RI. Figure 5-4 provides examples of aquatic ecological receptors that are preliminarily identified for further evaluation during the risk assessment based on the current understanding of the Site and consideration of the results of other CERCLA risk assessments performed at nearshore cleanup sites in the region. Exposure pathways relevant to these species are indicated on Figure 5-4 and include the following:

- Piscivorous Mammals (e.g., Harbor Seals). The potential for limited exposure
  exists for piscivorous mammals foraging at the Site. Potentially complete
  exposures are associated primarily with consumption of aquatic biota, and to a
  lesser extent with exposure to sediment and surface water.
- **Piscivorous Raptors (e.g., Ospreys).** The potential for limited exposure exists for piscivorous raptors foraging at the Site. Potentially complete exposures are associated primarily with consumption of aquatic biota, and to a lesser extent with exposure to surface water.
- Shore Birds (e.g., Herons and Sandpipers). The potential for exposure exists for shore birds residing or foraging at the Site. Potentially complete exposures are associated primarily with consumption of aquatic biota, incidental ingestion of sediment and to a lesser extent with exposure to surface water.
- **Piscivorous Fishes (e.g., Rockfish).** The potential for exposure to Site sediments and surface water exists for piscivorous fishes residing or foraging at the Site.
- Omnivorous Fishes (e.g., Sculpins). Omnivorous fishes residing or foraging at the Site may potentially be exposed to Site sediments and surface water.
- Benthivorous Fishes/Shellfish (e.g., Flatfish, Bivalves, and Crabs). Benthivorous fish/shellfish residing or foraging at the Site may potentially be exposed to Site sediments and surface water at the Site.
- Benthic Invertebrates (e.g., Benthic Infauna Community). Benthic invertebrates residing at the Site may potentially be exposed to site sediments and pore-water.
- Macrophytes (e.g., Algae and Kelp). Macrophytes residing at the Site may potentially be exposed to site sediment and surface water.

Data needed to refine the exposure pathways and receptors are identified in Section 9. These data needs include sampling and analysis of upland soils, groundwater, sediments, and biological receptors.

The upland properties at the Site have historically been developed and used for industrial operations. However, portions of these properties include habitat that could be used by terrestrial ecological receptors. These areas primarily include the vegetated areas of the Former Ravine and the bank and the beach. The risk assessment will include an evaluation of terrestrial receptors with differing modes of exposures such as nesting, foraging, residence, and/or presence at the Site. Representative receptors will be selected to evaluate the different exposure pathways. Preliminary incomplete terrestrial ecological receptors will be further evaluated as part of the RI. Figure 5-5 provides examples of terrestrial ecological receptors, which are preliminarily identified for further evaluation during the risk assessment based on the current understanding of the Site and consideration of the results of other CERCLA risk assessments performed at nearshore

cleanup sites in the region. Exposure pathways relevant to these species are indicated on Figure 5-5 and include the following:

- Avian Predators (e.g., Robins). The potential for exposure exists for avian
  predators foraging or nesting at the Site. Primary exposure pathways for these
  receptors include the consumption of soil invertebrates and incidental ingestion
  of Site soil.
- Carnivores (e.g., Coyotes). The potential for limited exposure exists for carnivores foraging at the Site. Primary exposure pathways for these receptors include the consumption of soil invertebrates and small mammals and incidental ingestion of Site soil.
- Omnivores (e.g., Raccoons). The potential for limited exposure exists for omnivores foraging at the Site. Primary exposure pathways for these receptors include the consumption of plants and soil invertebrates and incidental ingestion of Site soil.
- **Herbivores (e.g., Voles).** The potential for exposure exists for herbivores residing at the Site. Primary exposure pathways for these receptors include the consumption of plants and incidental ingestion of Site soil.
- Insectivores (e.g., Shrews). The potential for exposure exists for insectivores residing on the Site. Primary exposure pathways for these receptors include the consumption of soil invertebrates and incidental ingestion of Site soil.
- **Upland Vegetation.** The potential for exposure to Site soil exists for plants growing on the Site.
- **Soil Invertebrates.** The potential for exposure to Site soil exists for earthworms and other biota living in Site soil.

# 6 Project Planning

This section identifies initial potential ARARs, PRGs, and RAOs for the purposes of project planning. Potential ARARs are identified to facilitate communications with support agencies, help plan potential field activities, and assist in the identification of RAOs and PRGs. Initial PRGs are identified to help evaluate existing data and assist in the selection of appropriate analytical methods. ARARs, PRGs, and RAOs will be further developed during the RI/FS process. Those ARARs, PRGs, and RAOs that are determined to be applicable to the Site-related decisions may include some, none, or all of those identified in this section. The ARARs, PRGs, and RAOs that are ultimately determined to be applicable to the Site-related decisions will be established in consultation and coordination with key stakeholders and the public during the RI/FS process.

### 6.1 Applicable or Relevant and Appropriate Requirements

The project must comply with CERCLA Section 121, which requires remedial actions to achieve ARARs. According to the National Contingency Plan (Code of Federal Regulations, Title 40, Section 300.5 [40 CFR 300.5]), applicable requirements are those cleanup standards, standards of control, and other substantive requirements, criteria, or limitations promulgated under federal environmental or state environmental and facility siting laws that specifically address a hazardous substance, pollutant, contaminant, remedial action, location, or other circumstance identified at a CERCLA site. Appropriate and relevant requirements are cleanup standards, standards of control, and other substantive requirements, criteria, or limitations promulgated under federal environmental or state environmental or facility siting laws that are not applicable to a hazardous substance, pollutant, contaminant, remedial action, location or other circumstances at a CERCLA site, but address problems or situations similar to those encountered at the site that their use is well suited to the particular CERCLA site.

Some federal, state, and local environmental and health agencies may develop criteria, advisories, guidance documents, and proposed standards that are not legally enforceable but contain useful information for selecting cleanup levels or implementing a cleanup remedy. These fall into the category of "to be considered" (TBC) elements. TBCs are not mandatory requirements but may complement the identified ARARs.

ARARs and TBCs potentially relevant to the RI/FS are presented in Tables 6-1 through 6-3 and organized in the following categories:

- Contaminant-specific requirements;
- Location-specific requirements; and
- Performance, design, or other action-specific requirements.

Some ARARs fit neatly into a single category, while others may fall into more than one category. The categories are described as follows:

- Contaminant-specific ARARs are laws and requirements that establish health- or risk-based numerical values or methodologies for developing such values (EPA 1988b). These ARARs are used to establish the acceptable concentration of a contaminant that may remain in or be discharged to the environment. As such, contaminant-specific ARARs are considered in identifying the PRGs. Contaminantspecific ARARs are listed in Table 6-1.
- Location-specific ARARs are requirements that are triggered on the basis of the location of the remedial action to be undertaken (EPA 1988b). Location-specific ARARs may restrict or preclude certain remedial actions or may apply only to certain portions of the Site. Some location-specific ARARs overlap with actionspecific ARARs. Location-specific ARARs are listed in Table 6-2.
- Action-specific ARARs are performance, design, or other requirements that may place controls or restrictions on a particular remedial action (EPA 1988b). Action-specific ARARs are typically technology- or activity-based requirements or limitations on actions, and these requirements may include contaminant-specific standards or criteria that must be met as the result of an action. For remedial actions at the Site, these requirements are not necessarily triggered by the presence of specific contaminants in Site media, but rather by the specific actions that occur at the Site. Action-specific ARARs are listed in Table 6-3.

# 6.2 Remedial Action Objectives

RAOs consist of goals for protecting human health and the environment that are specific for each potentially contaminated environmental medium (e.g., soil, groundwater, and sediment). RAOs for protection of human receptors typically include both a contaminant level and an exposure route. RAOs for protection of environmental receptors typically seek to preserve or restore a resource and are typically expressed in terms of the medium of interest and target cleanup levels. The preliminary RAOs related to the protection of human health are as follows:

- Groundwater. Reduce risk to human health from direct contact with, and consumption of, groundwater contaminated with Site-related COCs to protective levels.
- **Sediment.** Reduce risk to human health from consumption of fish and shellfish containing Site-related COCs to protective levels.
- **Sediment.** Reduce risk to human health from incidental ingestion and/or dermal exposure to Site-related COCs during potential recreational use of the beach areas at the Site to protective levels.
- **Vapor.** Reduce risk to human health from inhalation of vapors from groundwater and/or soils contaminated with Site-related COCs to protective levels.

• **Soils (Surface and Subsurface).** Reduce risk to human health from direct contact with or incidental ingestion of Site-related COCs to protective levels.

The preliminary RAOs related to environmental protection are as follows:

- **Groundwater.** Reduce, to protective levels, risks to ecological receptors from direct contact with and consumption of groundwater contaminated with Siterelated COCs, including indirect exposure from consumption of prey exposed to groundwater entering the Port Washington Narrows.
- **Upland Soil.** Reduce, to protective levels, risks to terrestrial wildlife exposed to Site-related COCs through direct contact with and incidental ingestion of Site soil or consumption of soil-dwelling invertebrates.
- **Sediment.** Reduce, to protective levels, risks to aquatic wildlife from exposure to Site-related COCs in surface sediments or in prey species at the Site.
- **Sediment.** Reduce, to protective levels, risks to the benthos from Site-related COCs in surface sediments.

The preliminary RAOs will be developed further throughout the RI/FS process, in consultation with key stakeholders and the public, and may be revised, refined, or replaced.

# 6.3 Preliminary Remediation Goals

PRGs are published, generic, and conservative values that consider human health and ecological toxicity using standard exposure parameter values and risk assumptions to estimate protective chemical concentrations. Generic PRGs do not consider Site-specific conditions, exposure pathways, or potential receptors. An exceedance of a general PRG is not an indication of risk but an indication that further evaluation is required to determine risk. As additional information is collected throughout the RI/FS process, the PRGs will be modified to be directly applicable to Site conditions, exposure pathways, and receptors. This section identifies the initial PRGs for the screening of existing soil, groundwater, and sediment data. Initial surface water PRGs have been identified to assist with development of the RI/FS Work Plan; however, no surface water data are available for the Site.

Potential PRGs include numerical values identified in ARARs, peer-reviewed risk-based values, or values identified in other screening benchmark sources. Potential PRGs include values from the following sources:

#### ARARs:

- Soil: none available (except for those related to PCBs in the Toxic Substances Control Act);
- Groundwater: maximum contaminant levels (MCLs);

- Surface water: national recommended water quality criteria for human health (organism only) and aquatic life (chronic value); and
- Sediment: Washington State Sediment Management Standards (SMS).

#### Peer-reviewed sources:

- Soil: EPA human health regional screening levels (RSLs) and EPA ecological soil screening levels (EcoSSLs);
- Groundwater: EPA human health RSLs;
- Surface water: none available; and
- Sediment: NOAA effect range-low and effect-range-medium benchmarks (ER-L/ER-M) (Long et al. 1995).

### Other screening benchmark sources:

- Soil: EPA Region 5 Resource Conservation and Recovery Act (RCRA) EcoSSLs;
- Groundwater: none available;
- Surface water: EPA Region 3 Biological Technical Assistance Group (BTAG) sediment ecological screening benchmarks and EPA Region 5 RCRA ecological surface water screening levels; and
- Sediment: EPA Region 3 BTAG sediment ecological screening benchmarks and EPA Region 5 RCRA sediment ecological screening levels.

Tables 6-4, 6-5, 6-6, and 6-7 summarize the potential PRGs from these sources for each medium (soil, groundwater, sediment, and surface water, respectively) and identify an initial PRG for each contaminant. The initial PRG for a given contaminant was selected as the lowest of the ARARs or peer-reviewed risk-based criteria. If a value from these first two sources is unavailable, the initial PRG was selected as the lowest value in the "other screening benchmark" category. For sediment, the regionally specific SMS value was used. If no SMS value exists for the contaminant, the peer-reviewed NOAA value was used.

Identified initial PRGs include the following:

#### • Soil:

- EPA RSL residential,
- EPA RSL industrial,
- EPA EcoSSL birds,
- EPA EcoSSL mammals,
- EPA EcoSSL invertebrates,

- o EPA EcoSSL plants, and
- EPA Region 5 RCRA ecological screening levels for soil.

#### • Groundwater:

- o EPA MCL, and
- EPA RSL tap water.

#### Sediment:

- Washington State SMS sediment cleanup objective (SCO),
- NOAA ER-L benchmarks (Long et al. 1995),
- EPA Region 3 BTAG ecological marine sediment screening benchmarks, and
- o EPA Region 5 RCRA ecological sediment benchmarks.

#### • Surface water:

- o National recommended water quality criteria for aquatic life (EPA 2013a),
- EPA Region 3 BTAG ecological marine surface water screening benchmarks, and
- EPA Region 5 RCRA ecological surface water benchmarks.

For soil, two different initial PRGs were identified: one for surface soil (which includes consideration of screening levels for terrestrial ecological receptors) and one for subsurface soil at depths below potential ecological exposures.

# 7 Existing Data and Data Usability

Existing Site characterization data have been reviewed in terms of data usability for the RI/FS. The existing data include data for the Former Gas Works Property and also data for sediments and tissue within the Port Washington Narrows, Dyes Inlet, and nearby portions of Puget Sound.

# 7.1 Data Quality Characterization

Data quality review included the definition of minimum data acceptability criteria (MDAC). Relevant guidance was applied, including the following:

- EPA (1988a) Guidance for Conducting Remedial Investigations and Feasibility Studies Under CERCLA;
- EPA (1992) Guidance for Data Usability in Risk Assessment, Part A;
- EPA Contract Laboratory Program Function Guidelines for Data Review (variable dates for different analyte groups); and
- EPA (2009) Guidance for Labeling Externally Validated Laboratory Analytical Data for Superfund Use.

### 7.1.1 Minimum Data Acceptability Criteria

The MDAC evaluations of historical soil, groundwater, and sediment investigations in the ISA are described for each sampling event in Table 7-1. <sup>11</sup> MDAC evaluations of existing sediment and tissue data are described in Table 7-2. This MDAC review considered the following criteria:

- Work Plan Documentation:
  - Documentation describing the sampling program or event, the methods used, and the parties involved in sample collection must be available.
  - Collection methods must be clearly defined and be adequate for obtaining representative and quantitative information.
  - The purpose of data collection should be available.
- Sample Location and Collection Methods:

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<sup>&</sup>lt;sup>11</sup> Investigations conducted under the Order and performed in accordance with EPA-approved Quality Assurance Project Plans (QAPPs) (i.e., the 2013 TCRA) are not included in the MDAC tables.

- Sample coordinates and a qualitative understanding of accuracy (i.e., knowledge of how the location was established or the method by which the coordinates were obtained) must be documented. The coordinate system must be documented.
- Sample collection method and matrix must be documented. For example, a water sample must be identified as to whether it is a surface water, porewater, or groundwater sample and whether it is whole water or filtered (i.e., total versus dissolved fraction). Temporal or spatial compositing and sample volume must be identified. For tissue samples, tissue preparation must be documented.
- Sample depths and, where applicable, start and end depths must be identified.
- Sample storage methods must be documented and consistent with approved methods, including holding time and preservation.
- Sample chain of custody must be documented.

#### Laboratory Analysis:

- Data tables are available (not summary format) with laboratory reports and data validation information.
- Appropriate detection limits and quantitation limits are achieved so that the data meet the RI data quality objectives (DQOs) for environmental investigations:
  - Detection limits, units for each detection limit, and data qualifiers must be reported. Nondetected results must have the associated detection or reporting limits indicated. Data qualifiers must follow EPA guidance or be defined in documentation.
  - Analytical methods must be documented and acceptable based on EPA guidance.
  - Measurement instruments and calibration procedures must be documented.
  - Toxicity and bioaccumulation test methods must be documented, including any deviations from standard protocols. For risk assessment, test methods must follow standard protocols, including controls and reference tests. Proper documentation to assess methods and statistical treatment must be available. Where possible, statistical results should be recalculated from the raw test data.

- Taxonomic data must be reported to the lowest practicable taxonomic level on a sample-specific basis, with scientific nomenclature. Taxonomic levels must be sufficient to assess relevant metrics for ecological risk assessment, such as feeding guilds or stress-induced compositional changes in the community. Collection methods, sample preservation, and sample preparation methods must be documented.
- Biological community metric calculations must be defined and documented.
- Quality Control and Data Validation:
  - Documentation of field and laboratory quality control samples (duplicates, blanks) must be present.
  - Analytical chemical data must have been validated and qualified consistent with EPA functional guidelines or EPA Region 10 validation practices.
  - Hard copies of laboratory data reports (e.g., Form 1 or Certificates of Analysis) must be available to verify that electronic or tabulated data were accurately transcribed or transmitted.

### 7.1.2 Data Usability

Based on the results of the MDAC evaluation and considering the data representativeness for current Site conditions, the data were classified in one of the following data usability (DU) categories:

- **DU-1.** These data meet most or all of the MDAC requirements and are considered reasonably representative of Site conditions. DU-1 data are used in this Scoping Memorandum for COPC and source identification and preliminary evaluations of the nature and extent of contamination.
- **DU-2.** These data meet most of the MDAC requirements but have been superseded by more current or higher quality data for representation of the nature and extent of contamination. DU-2 data are used in this Scoping Memorandum for COPC and source identification.
- **DU-R.** These data do not meet the MDAC requirements and are not used in this Scoping Memorandum.

Of the existing data, the data were classified as follows:

- DU-1:
  - All data collected during the 2013 TCRA.

- Soil data, sediment data for analytes other than PAHs, and groundwater data from monitoring wells, collected during the 2008 TBA.
- Soil and groundwater data collected during the 2007 Preliminary Upland Investigation. These data met most of the MDAC criteria but underwent minimal data validation.
- Regional sediment monitoring data collected under the following programs:
  - 2008 and 2009 PSAMP Spatial/Temporal Monitoring, Central Sound
  - 1989 to 2013 PSAMP Long-Term/Temporal Monitoring
  - 2009 PSAMP Urban Waters Initiative, Bainbridge Basin
  - 2009 Ocean Survey Vessel Bold Summer 2008 Survey
- o 2010 and 2012 ENVVEST mussel data
- 2005 and 2007 NOAA Mussel Watch at station SIWP
- 2001 303d Ecology clam and crab sampling data

#### • DU-2:

 Sediment data collected during the 2010 TCRA and sediment data for PAHs collected during the 2008 TBA. These data met most of the MDAC criteria but have been superseded by more recent data collected in 2013, after the 2010 TCRA was completed.

#### • DU-R:

- Soil and sediment data collected during the 1995 Ecology Field Inspection.
  These data had limited documentation, including poorly documented
  sample locations, no documentation of collection or sample handling
  methods, and no chain of custody.
- Groundwater data collected from temporary borings during the 2008
   TBA. The samples were not filtered, and the data are not considered representative of groundwater conditions because of potential bias due to sample turbidity.

### 7.2 Existing Site-Related Data

This section summarizes the available data collected during previous investigations and removal actions conducted at the Site, relates that data to the CSM, and describes how the existing data might be used in the RI/FS. In this Scoping Memorandum, the existing data are used to develop a preliminary understanding of the nature and extent of contamination that will be further used in the RI/FS Work Plan to identify data gaps and

guide the Site investigation activities. Data identified in Section 7.1 as usable for this purpose, including data from the 2007 Preliminary Upland Investigation, selected data from the 2008 TBA, and data from the 2013 TRCA are presented below for Site media for which data are available (soil, groundwater, and sediment). Data classified as DU-1 (see Section 7.1) are included in the tables and figures in this section. Data summary tables for each medium that include all data classified as DU-1 or DU-2 are provided in Appendix D.

### 7.2.1 Soil Data

As discussed in Section 4.1, soil samples were collected as part of investigations conducted in 2007, 2008, and 2013. Soil samples were collected and analyzed for TPH, metals, SVOCs (including PAHs), VOCs, and PCBs. Table 7-3 summarizes the number of samples collected for analysis of each constituent and an evaluation of detected concentrations to the initial PRG. Data for metals are also compared to natural background concentrations. The soil analytical data are summarized in tables that are included in Appendix D.

The constituents detected in soil at concentrations above the initial PRGs include the following:

- VOCs, including benzene, ethylbenzene, cis-1,3-dichloropropene, and trans-1,3-dichloropropene;
- PAHs; and
- Metals, including antimony, arsenic, cadmium, chromium, cobalt, copper, lead, manganese, nickel, thallium, vanadium, and zinc.

Other than PAHs, no SVOCs were detected at concentrations above the initial PRGs; however, the reporting limits for a subset of SVOCs exceed the initial PRGs at some locations (Table 7-3 and Appendix D). Practical quantitation limits (PQLs) for COPCs will be identified in the RI/FS Work Plan to determine if lower detection limits are achievable or if the PRGs need to be adjusted.

PCBs were not detected in soil; the reporting limits for PCBs in all samples were less than the initial PRGs (Appendix D).

No initial PRGs were identified for TPH, which is not a CERCLA contaminant of concern. However, in the RI/FS, identifying the nature and extent of different TPH products (e.g., gasoline or diesel) may be helpful in defining contaminant sources. TPH data should be used with caution at sites, such as MGP sites, where non-petroleum hydrocarbon mixtures are present (e.g., coal tar). Therefore, an understanding of the type of product by chromatogram or other forensic analysis is needed to correctly interpret TPH data. For the purposes of this memorandum, TPH distribution was not evaluated but may be evaluated in the RI.

A summary of VOCs, PAHs, and metals detected at concentrations above the initial PRGs is provided in the following subsections by analyte group. The maximum concentration

detected at each boring location and a comparison to the initial PRGs and/or natural background concentrations in surface and subsurface soil is provided for the primary constituents detected at concentrations above the initial PRGs<sup>12</sup> (Figures 7-1 through 7-12). As described in Section 6.2, initial PRGs for surface soil include a consideration of potential terrestrial ecological exposure, whereas initial PRGs for subsurface soil do not. For the purposes of this preliminary evaluation, surface soil is defined as soils from 0 to 10 feet in depth, and subsurface soil is defined as 10 feet in depth or greater.

#### 7.2.1.1 VOCs

Two BTEX compounds, benzene and ethylbenzene, were detected at concentrations above the initial PRGs. The most frequent detections of benzene at concentrations above the initial PRG occurred at two locations: in surface soil collected at sample locations MW-3, in the vicinity of the former finished gas storage tanks, and SP03, near the edge of the Former Ravine fill area (Figure 7-1). Benzene was not detected in any subsurface soil samples at a concentration above the initial PRG (Figure 7-2).

Two halogenated VOCs, cis-1,3-dichloropropene and trans-1,3-dichloropropene, were detected at concentrations above the initial PRG in one sample. The source of these VOCs is unknown.

BTEX compounds are potentially an indicator of MGP-related releases but may result from other sources (e.g., gasoline-range TPH or industrial solvents). The existing data for BTEX in soil are used in this memorandum to help identify the upland ISA (see Section 8.2.1). The data will also be used in the RI to help assess the nature and extent of contamination. Additional data on the lateral and vertical extent of VOCs in soil are needed to evaluate potential source areas, delineate the extent of contamination, and determine risks to human health and the environment.

#### 7.2.1.2 PAHs

Figures 7-3 and 7-4 depict the maximum concentrations of naphthalene in surface and subsurface soil, respectively. Figures 7-5 and 7-6 depict the concentrations of total carcinogenic PAHs (cPAHs)<sup>13</sup> in surface and subsurface soil, respectively. The vertical distribution of naphthalene concentrations in soil is illustrated along geologic cross sections A–A′, B–B′, C–C′, and D–D′ in Figures 3-4 through 3-7, respectively.

The concentrations of total cPAHs and naphthalene exceeding the initial PRGs were detected at sample locations that correspond to operational areas of the former gas

<sup>&</sup>lt;sup>12</sup> Primary constituents shown on the figures include those detected with the greatest frequency or magnitude above the initial PRGs and natural background concentrations.

<sup>&</sup>lt;sup>13</sup> Concentrations of total cPAHs are provided in benzo(a)pyrene toxicity equivalent concentrations.

works. In surface soil, the highest concentrations of both total cPAHs and naphthalene were detected at sample location MW-3, advanced in the vicinity of the storage tanks, which held light oil and coal tar (Simonson 1997b). Likewise, the highest concentrations of both total cPAHs and naphthalene in subsurface soil were detected at sample location MW-6, which was advanced at the location of the former gas holder.

Generally, concentrations of naphthalene and cPAHs on the Former Gas Works Property are highest in surface soil and decrease with depth (MW-3 and SP03, for example). However, at MW-6, advanced at the location of the former gas holder, PAH concentrations detected in subsurface soil were much higher than those in surface soil. Because the gas holder was reportedly at least 10 feet deep, this finding may indicate that the gas holder was filled with cleaner soil after it was demolished. Also, the concentrations of PAHs detected in deeper soil were greater than those in shallow soil at well MW-8, located hydraulically downgradient of the former gas works operational area.

The concentrations of total cPAHs exceeding the initial PRG have been detected in soil samples collected between depths of 3 and 40 feet. The highest concentrations of total cPAHs were detected in shallow soil, between the depths of 5 and 12 feet, at well MW-3, well MW-6, and boring SP03 and in deeper soil at a depth of 25 feet at well MW-8.

The presence of cPAHs and naphthalenes is a potential indicator of MGP-related releases. <sup>14</sup> The existing data for PAHs in soil are used in this memorandum to help identify the upland ISA (see Section 8.2.1). The data will also be used in the RI to help assess the nature and extent of contamination. Additional data on the lateral and vertical extent of PAHs in soil are needed to evaluate potential source areas, delineate the extent of contamination, and assess risks to human health and the environment.

#### 7.2.1.3 Metals

The detectable concentrations or analytical reporting limits for a number of metals exceeded the initial PRGs. However, the concentrations of many of these metals did not exceed the natural background concentrations<sup>15</sup> (Ecology 1994):

 For manganese and antimony, all of the detected concentrations, and most of the reporting limits, are below the background concentrations.<sup>16</sup>

<sup>&</sup>lt;sup>14</sup> Carcinogenic PAHs and naphthalenes can also originate from other sources, including petroleum hydrocarbons or creosote. Forensic analyses, such as PAH fingerprinting, may be useful in the RI to help distinguish and identify potential sources of contamination.

<sup>&</sup>lt;sup>15</sup> Puget Sound background concentrations of metals were used for screening when available. When not available, Washington State background concentrations were used.

<sup>&</sup>lt;sup>16</sup> The Puget Sound regional background concentration for antimony has not been researched. The background concentration referenced is based on regional data from the Spokane Basin.

- Cobalt and vanadium were detected in all of the soil samples analyzed for metals, with many concentrations exceeding the initial PRGs; however, the detected concentrations are generally within the range of regional background concentrations.
- Thallium was detected at concentrations above the initial PRGs in most of the soil samples analyzed; a natural background concentration for thallium was not identified for this evaluation.

Detected concentrations of cadmium, lead, and zinc are within the range of regional background concentrations at most sample locations, except for borings MW-5, MW-8, and SP03, which are located at the northeast corner of the Former Gas Works Property in the shoreline and Former Ravine fill areas.

Arsenic, chromium, copper, and nickel were detected at concentrations above the initial PRGs and background concentrations at several locations. Figures 7-7 through 7-12 depict the concentrations of arsenic, copper, and nickel<sup>17</sup> in surface and subsurface soil. Concentrations of these metals in subsurface soil do not exceed the initial PRGs, with the exception of arsenic, which was detected at a concentration in excess of the initial PRG but less than the natural background concentration. Concentrations of arsenic, copper, and nickel in surface soil exceed the initial PRGs and the natural background concentrations at several locations. Arsenic was detected at concentrations above the natural background concentration at two locations: SP03 (Former Ravine fill area) and MW-3 (within the footprint of former gas works operations and the current industrial park). Copper, chromium, and nickel were sporadically detected across the Former Gas Works Property at concentrations above the natural background concentrations, and the maximum concentrations of copper, chromium, and nickel were only slightly greater than their respective background concentrations (62.7 milligrams per kilogram [mg/kg] versus 38 mg/kg for copper; 60.8 mg/kg versus 48 mg/kg for chromium; and 60.9 mg/kg versus 48 mg/kg for nickel). The sources of these exceedances are unclear from the existing data. Possible sources include contaminated fill, historical industrial operations, or natural background variability.

The existing soil data are useful for a preliminary identification of COPCs and provide an initial understanding of metals occurrences in surface and subsurface soil. These data can likely be used in the RI to inform the nature and extent of contamination. Additional data, particularly in surface soils and fill areas, are needed to evaluate potential sources

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<sup>&</sup>lt;sup>17</sup> Arsenic, copper, and nickel were mapped in soil because these constituents were also most frequently detected in groundwater at concentrations above the surface water or groundwater initial PRGs.

and delineate the extent of specific metals in soil, including arsenic, cadmium, chromium, copper, lead, nickel, and zinc.

### 7.2.2 Groundwater Data

As discussed in Section 4.1, groundwater samples were collected as part of the investigations conducted in 2007 and 2008. Groundwater samples were collected and analyzed for petroleum hydrocarbons, metals, SVOCs including PAHs, VOCs, and PCBs. Table 7-4 summarizes the number of samples collected for analysis of each constituent and the results of a comparison of detected concentrations to the screening criteria, which include concentrations protective of groundwater and surface water. The groundwater analytical data are provided in Appendix D.

The constituents detected in groundwater at concentrations above the initial PRGs include the following:

- **Metals:** arsenic, beryllium, chromium (both total and hexavalent), cobalt, copper, lead, manganese, nickel, thallium, vanadium, and zinc;
- **PAHs:** acenaphthene, benzo(g,h,i)perylene, dibenzofuran, phenanthrene, pyrene, naphthalenes, and total cPAHs;
- Pentachlorophenol (PCP); and
- **VOCs:** benzene, ethylbenzene, xylenes, 1,2,4-trimethylbenzene, 1,2-dichloroethane, carbon tetrachloride, chloroform, isopropylbenzene, n-hexane, and trichloroethene.

Other than the above-listed constituents, no SVOCs or VOCs were detected at concentrations above the initial PRGs; however, the reporting limits for a subset of SVOCs and VOCs exceed the initial PRGs at a number of locations (Table 7-4 and Appendix D). PCBs were not detected in groundwater; however, the reporting limits for PCBs in all samples were above the potential groundwater initial PRG (Appendix D).

The existing groundwater data are limited, with one sampling event at 10 locations and no groundwater data collected since 2008. The data are useful for the preliminary identification of COPCs, and they indicate where groundwater impacts may be located. These data can be used to support the development of the scope of work for the RI to evaluate the full lateral and vertical extent of COPCs in groundwater. The existing data, which were collected from wells that are still in place, can likely be used for future monitoring and may also be useful in the RI to evaluate long-term trends in groundwater quality.

VOCs, PAHs, PCP, and metals detected at concentrations above the initial PRGs are discussed in the following subsections by analyte group. The concentration detected at each monitoring well and a comparison to the groundwater initial PRGs are provided for

the primary constituents detected at concentrations above the initial PRGs<sup>18</sup> on Figures 7-13 through 7-17.

### 7.2.2.1 VOCs

One or more of the BTEX compounds were detected in groundwater samples collected at all of the monitoring wells except for wells MW-1 and SP02. The detected concentrations of benzene in groundwater are depicted on Figure 7-13. The highest concentrations were detected in wells MW-3, MW-6, and MW-8 (in and downgradient of the former gas works operation area).

#### 7.2.2.2 PAHs

Detected concentrations of total cPAHs were above the initial PRGs in groundwater samples collected from wells MW-3 through MW-8 (Figure 7-14) located on the Former Gas Works Property. The highest concentration of total cPAHs in groundwater was detected at well MW-4. There were no detected concentrations of cPAHs in the groundwater samples collected from wells MP04, SP02, MW-1, and MW-2.

The results for other PAHs are the following:

- Dibenzofuran and pyrene were detected at concentrations above the initial PRGs in the groundwater sample collected from well MW-4; and
- Naphthalenes, including 1-methylnaphthalene and naphthalene, were detected in groundwater samples collected from wells SP02, MP04, MW-3, MW-4, MW-5, MW-6, MW-7, and MW-8 at concentrations exceeding the initial PRGs. The highest concentrations of naphthalene were detected at wells MW-4 and MW-8 (Figure 7-15).

#### 7.2.2.3 Pentachlorophenol

PCP was detected in groundwater at a concentration exceeding the groundwater and surface water initial PRGs at well MW-8.

#### 7.2.2.4 Metals

The highest concentrations of metals in groundwater were generally detected at wells MW-3 and MW-4. MW-3 is located in the central portion of the Former Gas Works Property in the vicinity of the former finished gas storage tanks and former metal finishing operations. MW-4 is located within the Former Ravine fill area in the central portion of the Sesko Property. Results for specific metals are the following:

<sup>&</sup>lt;sup>18</sup> Primary constituents shown on the figures include those detected with the greatest frequency or magnitude above the groundwater initial PRGs.

- Arsenic was detected in all of the groundwater samples analyzed, at concentrations ranging from 0.6 to 26 micrograms per liter (μg/L), all of which exceed both the groundwater initial PRG and the surface water initial PRG. Figure 7-16 depicts the concentrations of arsenic in groundwater, which are highest in the central portion of the Former Gas Works Property, at wells MW-3 and MW-4.
- Hexavalent chromium was detected in groundwater samples collected from wells MW-1 and MW-3 through MW-8 at concentrations exceeding the groundwater initial PRG. The concentrations detected in wells MW-5 and MW-8 also exceed the surface water initial PRG. Figure 7-17 depicts the concentrations of hexavalent chromium in groundwater.
- Total chromium and lead were detected in groundwater at concentrations above both the groundwater initial PRGs and the surface water initial PRGs in the groundwater samples collected from wells MW-3 and MW-4.
- Copper and nickel were detected at concentrations exceeding surface water initial PRGs at most of the sample locations; none of the concentrations of copper and nickel exceeds the groundwater initial PRGs. The highest concentrations of copper and nickel were detected in groundwater samples collected from wells MW-3 and MW-4.
- Concentrations of cobalt, manganese, thallium, and vanadium exceeding the groundwater initial PRGs were detected in the groundwater sample collected from well MP04.

Potential sources of metals in groundwater include fill materials and historical industrial operations. More information is needed to determine the source and extent of metals in groundwater.

### 7.2.3 Sediment Data

Available sediment data for the Site include those collected in 2008 as part of the TBA, in 2010 as part of the 2010 TCRA, and in 2013 as part of the 2013 TCRA. These data sets include the following:

- **2008.** Five surface sediment samples from the beach north of the Former Gas Works Property were analyzed for TPH, VOCs, SVOCs, and metals.
- **2010.** Thirty-two surface sediment samples collected during the 2010 TCRA area were analyzed for VOC and SVOCs.
- **2013.** Thirty-nine surface sediment samples collected during the intertidal sediment sampling program were analyzed for total solids (TS), total organic carbon (TOC), and SVOCs.
- **2013.** Seventeen subsurface sediment samples were collected by direct-push methodology at seven locations. Samples from 4 discrete intervals were analyzed

for VOCs, and samples from 17 subsurface intervals were analyzed for TS, TOC, and SVOCs.

Table 7-5 presents these sediment data and the initial PRGs identified in Section 6.2. Where applicable, reference values are also presented for natural background concentrations of contaminants in Puget Sound sediments or soils.

Figures 7-18 through 7-22 present the measured concentrations of PAHs in beach sediments at the Site. Data are presented on a dry-weight basis for benzo(a)pyrene, total low-molecular-weight PAHs (LPAHs), total high-molecular-weight PAHs (HPAHs), total cPAHs, and total cPAH toxic equivalent (TEQ) concentrations. The highest PAH concentrations were detected within and near the two removal action areas. East and west of these two areas, concentrations decrease rapidly.

## 7.2.4 Surface Water Data (None)

No surface water data for the Former Gas Works Property or adjacent areas were identified as of the preparation of this Scoping Memorandum.

## 7.2.5 Tissue Data (None)

No tissue data for the Former Gas Works Property or adjacent areas were identified as of the preparation of this Scoping Memorandum.

## 7.3 Existing Data from Other Cleanup Sites

As described in Section 4.3.1, soil and groundwater data collected on the Former SC Fuels Property include TPH, BTEX, and lead. The majority of the soil data were collected prior to and during remedial actions (removal of USTs and surrounding contaminated soil), which occurred in 2002. The most recent groundwater monitoring data are from January 2007. During that sampling event, concentrations of benzene were detected in groundwater at concentrations up to 88  $\mu$ g/L on the Former SC Fuels Property and up to 49  $\mu$ g/L in the eastern portion of the Pennsylvania Avenue right-of-way (GeoScience Management 2007). The extent of benzene detected in groundwater (detection limit 1  $\mu$ g/L) in 2007 is shown on Figure 7-23.

## 7.4 Data for Port Washington Narrows and Dyes Inlet

A number of high-quality sediment and tissue studies were identified for the Port Washington Narrows and Dyes Inlet. The location of sediment and tissue data with measured PAH concentrations is shown on Figure 7-24. These data sets are not used for data screening or COPC evaluation (see Section 8.1) but provide valuable information about conditions in the vicinity of the Site.

## 7.4.1 Sediment Quality Data

Figures 7-25 and 7-26 present measured concentrations of benzo(a) pyrene and total cPAHs in sediments, respectively. Data are presented on a dry-weight basis. Ecology's current Draft Sediment Cleanup User's Manual II (Ecology 2013) recommends the use of the 90<sup>th</sup> percentile from data sets to evaluate natural and regional background concentrations. The 90<sup>th</sup> percentile concentrations of benzo(a) pyrene and total cPAHs in surface sediment samples collected during the Bold Survey in 2008 (USACE 2009) are approximately 10 micrograms per kilogram [µg/kg] and 50 µg/kg, respectively. Relative to the 90<sup>th</sup> percentile of the 2008 data, the concentrations of benzo(a) pyrene and total cPAHs in sediments from within the Port Washington Narrows, Dyes Inlet, and Sinclair Inlet are elevated. The vast majority of the measured values exceed the 90th percentile values from the 2008 data set.

The measured dry-weight concentrations of LPAHs and HPAHs in sediment are presented in Figures 7-27 and 7-28, respectively. The 90th percentile concentrations of LPAHs and HPAHs in surface sediment samples collected during the 2008 Bold Survey are 10.9  $\mu g/kg$  and 75.1  $\mu g/kg$ , respectively. Relative to the 90th percentile of the 2008 data, the LPAH and HPAH concentrations measured in Port Washington Narrows, Dyes Inlet, and Sinclair Inlet show the same magnitude of elevated concentrations as that shown in the cPAH data.

Existing sediment data sets may be used during the RI/FS to document existing sediment quality within nearby portions of the Port Washington Narrows and Dyes Inlet. The data may be useful, along with the data collected during the RI/FS, in evaluating the recontamination potential for Site sediments.

## 7.4.2 Tissue Quality Data

Figures 7-29 and 7-30 provide a synopsis of available existing PAH testing data for various aquatic organisms. Tested organisms include mussels, clams, and crabs. The data for total cPAHs are presented on both a wet-weight basis (Figure 7-29) and a lipid-normalized basis (Figure 7-30), respectively.

These tissue data sets may be useful during the RI/FS for evaluating how contaminant levels in tissues at the Site (predicted or empirically measured) compare to those in other seafood collected within the region.

## 7.4.3 Water Quality Data

No current water quality data for chemical contaminants within the Port Washington Narrows have been identified as of the preparation of this Scoping Memorandum.

Several studies have been conducted to assess potential contaminant inputs to Dyes Inlet and adjacent waters (Crecelius et al. 2003). The results of these and other available studies may be used qualitatively for the evaluation of potential nonpoint sources of pollution but will not be relied upon for the baseline risk assessment.

# 8 RI/FS Approach

### 8.1 Contaminants of Potential Concern

This section identifies preliminary COPCs based on: (1) contaminants typically associated with the former gas works process (carbureted water gas); (2) contaminants associated with other potential historical sources within the ISA (see Section 8.2); (3) contaminants detected during previous Site investigations; and (4) other EPA contaminants of interest. The COPCs, and ultimately the COCs, that are determined to apply to the Site-related decisions may include some, none, or all of the contaminants identified in this section. The COCs that are ultimately determined to apply to the Site-related decisions will be established on the basis of data and information that is collected as part of the RI/FS process.

Contaminants typically associated with carbureted water-gas manufacturing processes include the following:

- Light aromatic hydrocarbons, such as BTEX compounds;
- Heavier aromatic hydrocarbons, including PAHs;
- Other SVOCs, such as tar acids (e.g., phenol and cresols) and heterocyclic aromatics (e.g., carbazole and dibenzofuran); and
- Cyanide and sulfides associated with spent purifier materials.

COCs identified at a number of other nationwide MGP sites<sup>19</sup> are summarized in Table 8-1. COCs typically associated with MGPs include PAHs, BTEX, and cyanide.

Other historical processes with the potential for releases within the ISA include petroleum transfer and storage, metal fabrication, and vehicle and equipment salvage and repair. Contaminants typically associated with these processes include solvents (VOCs), petroleum hydrocarbons (including BTEX and PAHs), and metals.

Available Site data for soil, groundwater, and sediment are compared to the initial PRGs in Section 7. Contaminants detected at concentrations above the initial PRGs (and natural background concentrations, for naturally occurring metals) include the following:

VOCs, including benzene, ethylbenzene, xylenes, 1,2,4-trimethylbenzene, cis-1,3-dichloropropene, trans-1,3-dichloropropene,1,2-dichloroethane, carbon tetrachloride, chloroform, and trichloroethene;

<sup>&</sup>lt;sup>19</sup> Table 8-1 includes representative nationwide MGP sites at which the site conditions are similar and for which cleanup is in progress or has been completed (see Section 8.5).

- SVOCs, including PAHs and PCP; and
- Metals, including antimony, arsenic, cadmium, chromium, cobalt, copper, lead, manganese, nickel, thallium, vanadium, and zinc.

Other EPA contaminants of interest consist of polychlorinated biphenyls (PCBs) and pesticides. PCBs are man-made organic chemicals, manufactured between 1929 and 1979, and used in industrial and commercial applications including electrical, heat transfer, and hydraulic equipment; in paints, plastics and rubber products; and in pigments and dyes. PCBs may still be present in products and materials that were manufactured before 1979, including electrical transformers and capacitors, fluorescent light ballasts, adhesives, oil-based paint and caulking. Pesticides are substances, or mixtures of substances, intended for preventing, destroying, repelling, or mitigating any living organisms (e.g. insects, mice, weeds, fungi, microorganisms) that occur where they are not wanted or that cause damage to crops, humans or other animals. The term pesticide applies to insecticides, herbicides, fungicides, and various other substances used to control pests.

Table 8-2 provides a summary of the preliminary Site COPCs and includes the basis for their inclusion and why they are a concern. The preliminary Site COPCs include the following contaminant groups:

- VOCs, as identified and quantified by EPA Method 8260C.
- SVOCs, including carcinogenic- and non-carcinogenic PAHs, as identified and quantified by EPA Method 8270D/SIM.
- Metals, as identified and quantified by EPA Methods 200.8/6010/6020/7471B.
- PCBs, as identified and quantified by EPA Method 8082.
- Pesticides, as identified and quantified by EPA Method 8081B.
- Cyanide, as identified and quantified by EPA Method 9014.

Specific contaminants are listed on Table 8-2, by contaminant group, if information indicates they are confirmed or suspected to be present at the Site. However, the list of specific contaminants on Table 8-2 is not intended to be an exhaustive and complete list of preliminary Site COPCs. The scope of work for the RI/FS will include collection and analysis of samples from each media for the full standard list of contaminants for each contaminant group. Throughout the RI/FS process, the list of preliminary Site COPCs will be evaluated and revised as data is collected.

## 8.2 Initial Study Area

As described in the Statement of Work (SOW) for the AOC, the purpose of the ISA is to focus sampling and analysis in the first phase of the RI/FS. <sup>20</sup> The ISA is not intended to define the Site boundaries. <sup>21</sup>

The SOW anticipates "the ISA will encompass the area of operation of a former manufactured gas plant (MGP)..., including the area where contaminants from the area of operation have come to be located, which includes upland, beach and sediments." The ISA has been developed following the guidelines established by the SOW. The rationale for the ISA is further explained in Sections 8.2.1 and 8.2.2.

## 8.2.1 Upland Portion of Initial Study Area

The upland portion of the ISA (Figure 8-1) includes the Former Gas Works Property and portions of neighboring properties where gas works operations, including byproduct storage and disposal, are documented or suspected to have occurred. The upland portion of the ISA also includes the northern portion of the Penn Plaza Property where a drip tank was located and the other portion of the Sesko Property where materials from the former gas works process may have been placed in the Former Ravine. The upland portion of the ISA also includes areas where contamination not associated with the former gas works could potentially be commingled with gas works contamination. These non-gas-works operations include the former Lent's bulk petroleum storage tank farm on the Sesko Property, petroleum pipelines located in the northern portion of the Penn Plaza Property and the Sesko Property, and various light industrial operations on the McConkey and Penn Plaza Properties.

Consistent with the SOW, the proposed ISA encompasses all upland areas where contaminants associated with the former gas works are likely to be located. The existing data collected from areas near the boundaries of the ISA suggest that contamination associated with the former gas works may not extend beyond the ISA. More data are needed to determine if this is the case. The existing data include the results of soil and groundwater sampling from well MW-1 on the Penn Plaza Property, borings MP03 and MP02 within Thompson Drive, borings SP01 and SP02 on the Sesko Property, and explorations associated with the Former SC Fuels Property to the east of the ISA.

The first phase of the RI will characterize the nature and extent of contamination within the ISA and assess the subsurface characteristics that may influence the migration of contaminants. These data will be used to determine where additional investigation may

<sup>&</sup>lt;sup>20</sup> SOW, Sections 1.1 and 3.1.11.

<sup>&</sup>lt;sup>21</sup> SOW, Section 1.1.

be warranted. Investigations outside of the ISA, if needed, would then be specifically designed and implemented to focus on characterization of identified issues.

## 8.2.2 Sediment Portion of Initial Study Area

The sediment portion of the proposed ISA (Figure 8-2) comprises intertidal and subtidal areas in the general vicinity of the Former Gas Works Property. The sediment ISA is described as follows:

- Historical potential source areas associated with the former gas works (including the Former Gas Works Dock and the former drainage line) have been included.
- All beach sediments adjacent to the Former Gas Works Property that exhibited elevated PAH concentrations during the 2013 TCRA have been included.
- The offshore boundary of the ISA extends out past midchannel in the Port
  Washington Narrows, well past the bathymetric low point in the channel. This
  addresses potential migration pathways associated with groundwater and/or
  NAPL migration and those associated with potential sediment transport.
- The eastern and western boundaries of the ISA extend between 500 and 1,000 feet in an east-west direction from the Former Gas Works Property, allowing documentation of the potential transport of sediments that may have resulted from the east-west tidal currents occurring within the Port Washington Narrows.

The ISA includes multiple potential sources that are unassociated with historical activities on the Former Gas Works Property: multiple historical petroleum transfer docks, multiple stormwater and CSO outfalls, and the Port Washington Marina.

As part of the RI/FS activities related to sediments, there is a need to understand trends in sediment quality or water quality that may affect either current Site conditions or could potentially result in future recontamination of the Site. Therefore, sampling activities for sediments and surface water will not be exclusively confined to the ISA. Some sampling during the RI/FS will occur outside the sediment portion of the ISA.

## 8.3 Investigation Methods

Implementation of numerous investigation methods may be appropriate to fill the identified data gaps. The methods discussed herein are general approaches that will be considered for use during the RI. The RI/FS Work Plan will present the specific details of the investigation methods and approaches for the RI. The methods will include those that are appropriate to address the specific data needs and have been tested and demonstrated to be effective at similar sites with similar physical characteristics. Previous investigations in the ISA have included hollow-stem auger borings to collect soil samples and install wells to depths of 45 feet. Direct-push soil borings have been used for soil sampling in the upper 16 feet at the adjacent Former SC Fuels Property. A limited-access direct-push drilling rig encountered impenetrable native sediments at depths of 3 to 4 feet. Advancing into the dense native soils beneath the shallow fill material with the

use of direct-push drilling methods will likely be difficult. Impenetrable soil due to debris (e.g., wood or concrete) may also be encountered in fill areas such as the shoreline or the Former Ravine.

Fill soil and shallow native soils will likely be best characterized by means of a combination of exploration excavations (i.e., test pits or trenches using a backhoe or excavator) and direct-push soil borings where excavator access is limited (e.g., beneath buildings). Deeper native soils, in which soil borings are likely to be less effective at reaching the targeted exploration depths, will likely be best characterized by means of drilling methods that use heavier hammers and larger diameter augers (e.g. hollow-stem augers or sonic drills).

The methods for evaluating the presence and degree of contamination will include visual observation and chemical analytical results. Therefore, the collection of sufficient soil samples by means of competent drilling methods will be crucial to the success of the investigation. Likewise, properly constructed and developed monitoring wells will be necessary. Given the observed depth to groundwater during previous investigations, the wells can likely be installed using hollow-stem auger or sonic drilling methods. To minimize carrydown, use of a double-cased drill may be prudent for multilevel well installation in contaminated areas. The evaluation of groundwater flow, groundwater-surface water interaction, near-shore transition zone water, and migration of contaminants in groundwater will be performed with the use of a combination of investigation methods, which may include slug testing at upland monitoring wells and tidal studies.

The risk of vapor intrusion associated with volatile contaminants will be assessed using shallow soil, groundwater, and/or soil gas data. Soil gas may be evaluated using direct-push drilling methods to install shallow, temporary soil gas sampling points.

To meet the specific objectives, additional methods of assessing the presence, nature, and extent of contamination may be considered as the investigation activities progress. For example, the TarGOST® technology, which uses laser-induced fluorescence to delineate coal tar or creosote NAPL, could possibly be used to detect and characterize the extent of NAPL in fill and shallow native soils in areas where coal tar or creosote have been identified by other investigation methods. However, TarGOST® is specifically intended for use in delineating NAPL-contaminated zones and is appropriate only for sites where there is a confirmed presence of coal tar or creosote NAPL. A preliminary understanding of NAPL presence and occurrence in shallow or deeper soils would be needed to determine whether the use of TarGOST® would provide an advantage over more conventional exploration technologies. The use of electrical resistivity imaging may also be tool that could provide information about subsurface conditions at the Site. These and other assessment tools will be evaluated and potentially used during the RI/FS investigation.

## 8.4 Risk Assessment Methodology

Consistent with the AOC, a baseline ecological risk assessment (ERA) and a human health risk assessment (HHRA) will be performed to support RI/FS decision-making. The baseline risk assessments will be completed in parallel with the Draft RI Report. The RI/FS Work Plan will include additional details regarding the development of the ERA and HHRA. This section provides an overview of the exposure scenarios likely to be evaluated and the data needed to support those evaluations.

The preliminary CSM (Section 5) describes potentially complete exposure scenarios and pathways for human and ecological receptors. During the RI fieldwork, empirical data will be collected to quantitatively evaluate the level of risk for each receptor listed on Figures 5-3, 5-4, and 5-5. The needs related to risk assessment data, including both planned and contingent data collection needs, are identified in Sections 8.4.1 and 8.4.2.

#### 8.4.1 Human Health Risk Assessment

HHRA methodology will be based on national and regional guidance designated by EPA, including, but not limited, to the following:

- Risk Assessment Guidance for Superfund (RAGS), Volume I Human Health Evaluation Manual (Parts A through F);
- Interim Guidance: Developing Risk Based Clean-up Levels at Resource Conservation and Recovery Act Sites in Region 10 (January 1998);
- The 2011 Exposure Factors Handbook; and
- The 2007 Framework for Selecting and Using Tribal Fish and Shellfish
  Consumption Rates for Risk-Based Decision Making at CERCLA and RCRA Cleanup
  Sites in Puget Sound and the Strait of Georgia.

Toxicity data will be developed based on the EPA (2003) hierarchy of human health toxicity values.

Human exposure scenarios will be evaluated in the risk assessment for both beach/aquatic and upland areas of the Site. Scenarios to be evaluated for the beach/aquatic areas include the following:

- Recreational Beach Use. The potential for limited recreational beach use exists
  for individuals residing in proximity to the Site. Potential exposures to Site soil,
  sediment, and surface water will be addressed under this scenario.
- Fish/Crab Collection for Consumption. The portions of the Port Washington Narrows adjacent to the Former Gas Works Property currently support the collection and consumption of fish and crabs under WDFW regulations. In addition to the consumption of fish and crabs, potential exposure to Site sediment and surface water will be addressed under this scenario.

- Shellfish Collection for Consumption. The portions of the Port Washington Narrows adjacent to the Former Gas Works Property are currently listed as closed to shellfish harvesting (due to water quality concerns associated with CSOs and non-Site-related concerns) by the Washington State Department of Health; however, exposures associated with shellfish harvesting will be evaluated to understand potential risks should the shellfish harvesting restrictions be lifted. In addition to the consumption of shellfish, potential exposure to Site sediment and surface water will be addressed under this scenario.
- Beach Construction/Excavation Worker. Workers performing utility upgrades or maintenance or other activities may disturb sediments in the beach areas adjacent to the Former Gas Works Property. The potential risks resulting from exposures to Site surface and subsurface sediment and fugitive dust and vapor will be addressed under this scenario.

Human health risks associated with the upland areas of the Site will be evaluated as follows:

- Occupational Worker. The McConkey and Sesko Properties and the properties in the vicinity are zoned for industrial uses. The potential for limited exposures to Site surface soil and fugitive dust and vapor will be addressed under this scenario.
- Upland Construction/Excavation Worker. Workers performing utility upgrades or maintenance or other activities may disturb soils at the Site. The potential risks resulting from exposures to Site surface and subsurface soil and fugitive dust and vapor will be addressed under this scenario.
- Residential. The McConkey and Sesko Properties and the properties in the
  vicinity are zoned for industrial uses. However, exposures to residents will be
  evaluated to understand potential implications should these properties be
  converted to residential uses. The potential for limited exposures to Site surface
  soil and fugitive dust and vapor will be addressed under this scenario. Although
  no water supply wells are located on or near the former gas works, consumption
  of groundwater is retained as a pathway for screening, pending further
  evaluation of groundwater beneficial uses.

Data needed to support the HHRA, including both planned and contingent data collection needs, are the following.

- Definition of the nature and extent of Site-associated COCs in soils, groundwater, and sediment and potential concentrations of Site-associated COCs in surface water. Further testing and data screening are necessary to finalize the list of Siteassociated COCs in these media, as necessary to quantify exposure estimates.
- Information regarding potential seafood resources available at and near the Former Gas Works Property. This information is needed to better support the

development of exposure estimates related to the human consumption of seafood. This information includes further compilation of fish and shellfish abundance in the Port Washington Narrows and Dyes Inlet, video surveys of submerged areas within the ISA, and shellfish abundance surveys in beach areas at and near the Former Gas Works Property.

• Estimation of the potential concentration of Site-associated COCs accumulating in seafood at and near the Former Gas Works Property. This evaluation will initially be performed using bulk sediment, porewater, and surface water COC concentrations and bioaccumulation estimates derived from previous studies in the literature. If necessary, Site-specific tissue samples may be collected from selected species to validate and refine the initial estimates. The potential need for this contingent tissue sampling will be evaluated in coordination with EPA. If sampling is determined to be warranted, the methods will be documented in an RI/FS Work Plan Addendum.

Upland risk estimates associated with air quality (dust and vapors) will be initially developed using soil and groundwater data and model-derived estimates of dust and vapor concentrations. If necessary, collection of Site-specific soil vapor data may be conducted. The potential need for this contingent sampling will be evaluated in coordination with EPA. If sampling is determined to be warranted, the methods will be documented in an RI/FS Work Plan Addendum.

## 8.4.2 Ecological Risk Assessment

ERA methodology will address both terrestrial and aquatic ecological exposures. ERA methodology will be based on EPA guidance, including, but not limited to, the following:

- Ecological Risk Assessment for Superfund: Process for Designing and Conducting Ecological Risk Assessments, Interim Final, June 1997;
- Guidelines for Ecological Risk Assessment, 1998; relevant and appropriate updated EPA guidance material (e.g., EPA's Eco Updates); and
- EPA Region 10 Supplemental Ecological Risk Assessment Guidance for Superfund, 1997.

Toxicity data will be developed according to EPA guidance (e.g., EcoSSLs) and databases (e.g., ECOTOX), peer-reviewed scientific literature, and recent EPA-approved risk assessments.

Ecological exposure scenarios will be evaluated in the risk assessment for both aquatic and terrestrial ecological exposures. Terrestrial exposures to be evaluated include the following:

- Avian Predator (e.g., Robins). The potential for exposure exists for individuals foraging or nesting on the Site. Potential exposures to Site soil, terrestrial biota, and on-site water will be addressed under this scenario.
- Carnivore (e.g., Coyotes). The potential for limited exposure exists for individuals
  foraging on the Site. Potential exposures to Site soil, terrestrial biota, and on-site
  water will be addressed under this scenario.
- Omnivore (e.g., Raccoons). The potential for limited exposure exists for individuals foraging on the Site. Potential exposures to Site soil, plants and terrestrial biota, and on-site water will be addressed under this scenario.
- Herbivore (e.g., Voles). The potential for exposure exists for individuals residing
  on the Site. Potential exposures to Site soil, plants, and on-site water will be
  addressed under this scenario.
- Insectivore (e.g., Shrews). The potential for exposure exists for individuals residing on the Site. Potential exposures to Site soil, terrestrial biota, and on-site water will be addressed under this scenario.
- **Upland Vegetation.** The potential for exposure exists for plants growing on the Site. Potential exposures to Site soil will be addressed under this scenario.
- Soil Invertebrate. The potential for exposure exists for earthworms and other biota living in Site soil and will be addressed under this scenario.

The aquatic-dependent ecological exposure scenarios and key assumptions will include the following:

- **Piscivorous Mammal (e.g., Harbor Seals).** The potential for limited exposure exists for individuals foraging at the Site. Potential exposures to Site sediment, surface water, and aquatic biota will be addressed under this scenario.
- **Piscivorous Raptor (e.g., Ospreys).** The potential for limited exposure exists for individuals foraging at the Site. Potential exposures to Site surface water and aquatic biota will be addressed under this scenario.
- Shore Birds (e.g., Herons and Sandpipers). The potential for exposure exists for individuals residing or foraging at the Site. Potential exposures to Site sediment, surface water, and aquatic biota will be addressed under this scenario
- Piscivorous Fishes (e.g., Rockfish). The potential for exposure exists for individuals residing or foraging at the Site. Potential exposures to Site sediment porewater, surface water, and aquatic biota will be addressed under this scenario.
- Omnivorous Fishes (e.g., Sculpins). The potential for exposure exists for individuals residing or foraging at the Site. Potential exposures to Site sediment

porewater, surface water, and aquatic biota will be addressed under this scenario.

- Benthivorous Fishes/Shellfish (e.g., Flatfish, Bivalves, and Crabs). The potential
  for exposure exists for individuals residing or foraging at the Site. Potential
  exposures to Site sediment, porewater, surface water, and aquatic biota will be
  addressed under this scenario.
- Benthic Invertebrates (e.g., Benthic Infauna Community). The potential for exposure exists for individuals residing at the Site. Potential exposures to Site sediment and sediment porewater will be addressed under this scenario.
- Macrophytes (e.g., Algae and Kelp). The potential for exposure exists for individuals residing at the Site. Potential exposures to Site sediment, sediment porewater, and surface water will be addressed under this scenario.

Toxicity data will be developed according to EPA guidance (e.g., EcoSSLs) and databases (e.g., Ecotox), peer-reviewed scientific literature, and recent EPA-approved risk assessments.

Data needed to support the ERA, including both planned and contingent data needs, are the following:

- Definition of the nature and extent of Site-associated COCs in soils and sediment and potential concentrations of Site-associated COCs in surface water. Further testing and data screening are necessary to finalize the list of Site-associated COCs in these media, as necessary to quantify exposure estimates.
- Information regarding potential fish and wildlife resources available at and near
  the former gas works. This information is needed to better support the
  development of exposure estimates for the ERA. This information includes
  further compilation of fish and shellfish abundance in the Port Washington
  Narrows and Dyes Inlet, video surveys of submerged areas within the ISA, and
  shellfish abundance surveys in beach areas at and near the former gas works.
- Estimation of the potential concentration of Site-associated COCs accumulating in aquatic organisms at and near the former gas works. This evaluation will initially be performed using bulk sediment, porewater, and surface water COC concentrations and bioaccumulation estimates derived from previous studies in the literature. If necessary, Site-specific tissue samples may be collected from selected species to validate and refine the initial estimates. The potential need for this contingent tissue sampling will be evaluated in coordination with EPA. If sampling is determined to be warranted, the methods will be documented in an RI/FS Work Plan Addendum.

## 8.5 Potential Remedial Approaches

An understanding of potential remedial approaches that may be implemented at the Site is helpful during the scoping process to begin identifying data gaps, particularly for data needed to evaluate particular remedial technologies. Data gaps related to remedial technologies principally include site characterization data but may include bench- or pilot-testing of potential technologies if a need is identified during the RI/FS process.

This section describes potential remedial technologies and identifies remedial approaches that have been used at similar sites. Specific data needs for developing and evaluating potential remedial approaches will be described in the RI/FS Work Plan.

## 8.5.1 Remedial Technologies

Site remediation to achieve RAOs typically occurs by implementation of a combination of remedial technologies. Depending on the Site-specific circumstances, the use of remedial technologies may result in the complete elimination or destruction of hazardous substances at the Site, the reduction or elimination of migrating hazardous substances at the Site, or some combination of these effects. These technologies may be used in combination with engineering controls (e.g., barriers such as fences or caps) or institutional controls (i.e., non-engineered controls such as land use restrictions) when hazardous wastes remain at the Site. Remedial technologies are often categorized by the following general response actions:

- Monitored Natural Attenuation. Natural attenuation is the reduction of
  contaminant concentrations at the point of exposure over time by means of
  natural processes, such as sedimentation, sorption, dispersion, and/or
  biodegradation. Monitoring documents that the processes are occurring at the
  desired rates. For sediment, this general response action is referred to as
  monitored natural recovery.
- In Situ Containment. In situ containment involves confining hazardous substances in place by the placement of physical barriers or hydraulic controls. Containment technologies can be designed to prevent contact with and/or migration of hazardous substances.
- *In Situ* Treatment. *In situ* treatment technologies can potentially reduce the concentration, mobility, and/or toxicity of COCs.
- Removal. Contaminated materials can be physically removed from the Site and treated and/or disposed of at either an on-site or an off-site permitted disposal facility.
- **Ex Situ Treatment.** Ex situ treatment technologies destroy or immobilize contaminants in media that have been removed from the subsurface.

• **Disposal.** Disposal technologies include the placement of contaminated solid media in on-site or off-site landfills or the discharge of contaminated water to a publicly owned treatment works.

Preliminary lists of potential remedial technologies for NAPL, soil, groundwater, and sediment at the Site are provided in Tables 8-3 through 8-6.

## 8.5.2 Remedial Approaches at Other MGP Sites

Hundreds of MGP sites around the country have been through or are undergoing an RI/FS and cleanup action. Table 8-1 identifies remedial approaches that have been fully or partially implemented at MGP sites with characteristics (e.g., geology and presence of adjacent surface water bodies) that are similar to the Bremerton Gas Works Site. Common actions have included combinations of removal with off-site disposal or on-site treatment, solidification/stabilization, and institutional and engineering controls. Other technologies have included pump-and-treat, bioremediation, *in situ* chemical oxidation, barriers, and NAPL collection.

## 9 Summary and Data Gaps

Tables 9-1 and 9-2 summarize the principal data needs for the RI/FS that were defined during the initial scoping process.

Table 9-1 presents the data needs relating to the upland areas of the Site, including the data needed to support the risk assessment and FS activities for these areas. Table 9-2 presents the data needs for the beach and aquatic areas of the Site.

Most data gaps are to be filled during a single phase of field investigations. Potential investigation methods are discussed in Section 8.3. Specific proposed sampling methods and target locations will be defined in the RI/FS Work Plan. The anticipated sequence of field activities for upland and sediment areas will be defined in the RI/FS Work Plan but is expected to include the following:

#### Upland investigations:

- Complete ground-penetrating radar and utility locating.
- Conduct sampling of soils and fill material using direct-push borings, angled borings, test pits, trenches, and hand augers.
- O Characterize deep lithology and soil quality using deep borings.
- Complete selected borings as monitoring wells.
- Characterize Site hydrogeology, including performance of slug tests and a tidal study.
- Conduct quarterly groundwater monitoring.

#### Sediment investigation sequencing:

- Conduct video surveys to identify substrate, habitat characteristics, and presence/abundance of aquatic resources near the Site.
- Conduct beach surveys to evaluate the distribution of shellfish and other resources within and near the beach areas adjacent to the Former Gas Works Property.
- Sample and analyze surface sediments within the ISA to define the nature and extent of Site-related COCs. A subset of samples will be analyzed for PAHs in porewater to evaluate bioavailability of these contaminants.
- Sample and analyze surface sediments at selected locations beyond the ISA to supplement available data regarding sediment quality and potential recontamination sources within the Port Washington Narrows.

- Sample and analyze surface water at selected Site and background locations, including multiple sampling events to assess potential variability in surface water concentrations.
- Collect subsurface sediment core samples from the beach and subtidal areas sloping down into the Port Washington Narrows to evaluate the vertical distribution of Site-related COCs (including the potential presence of NAPL and hydrocarbon sheen) in subsurface sediments.
- Monitor near-bottom tidal currents within aquatic areas of the Site to assist in the evaluation of sediment stability.

After completion of the initial field program and consultation with EPA, some additional work may or may not be required to address contingent activities or to fully define the nature and extent of contamination at the Site. If applicable, these contingent or follow-up activities will be defined in an RI/FS Work Plan Addendum. Examples of work that might be defined as part of the RI/FS Work Plan Addendum include the following:

- Potential "step-out" sampling in the upland or sediment areas of the Site (if needed);
- Contingent sediment bioassay and/or seafood tissue testing if determined necessary for completion of the risk assessment; and
- Contingent sediment geochronology testing if determined necessary to support the evaluation of sediment stability and recovery processes.

Completion of treatability testing is not expected to be required to support the FS. However, this potential need will also be revisited after completion of the initial field program.

Preparation of the RI, risk assessment, and FS reports will be conducted in a manner that is consistent with the schedule requirements in the AOC.

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# **TABLES**

#### Table 3-1 - Monitoring Well Construction Information and Groundwater Elevation Measurements

Bremerton Gas Works Site Bremerton, Washington

							Depth to Water (feet below TOC)	Groundwater Elevation (feet NAVD88)
Well Identification	Installed By	Date Installed	Surface Elevation (Datum Unknown)	Total Boring Depth (Feet)	Depth to Top of Screen (Feet)	Depth to Bottom of Screen (Feet)	1-Jun-07	1-Jun-07
MP-04	E&E	5/13/2008	12.38	40	30	40		-
SP-02	E&E	5/12/2008	10.44	35	25	35		-
			Surface Elevation in feet (NAVD88)					
MW-1	GeoEngineers	5/21/2007	45.03	46.5	30	45	34.68	10.35
MW-2	GeoEngineers	5/21/2007	42.54	46.5	30	45	35.25	7.29
MW-3	GeoEngineers	5/22/2007	39.1	46.5	30	45	32.9	6.2
MW-4	GeoEngineers	5/23/2007	35.2	41.5	20	40	29.32	5.88
MW-5	GeoEngineers	5/24/2007	18.51	21.5	5	20	15.21	3.3
MW-6	GeoEngineers	5/22/2007	34.95	36.5	15	35	30.2	4.75
MW-7	GeoEngineers	5/23/2007	33.24	36.5	15	35	30.21	3.03
MW-8	GeoEngineers	5/22/2007	35.56	41.5	20	40	32.64	2.92

Notes:

-- = not measured

E&E = Ecology and Environment

NAVD88 = North American Veritcal Datum of 1988

TOC = top of casing

# Table 6-1 – Potential ARARs, Contaminant-Specific

Bremerton Gas Works Site Bremerton, Washington

Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness
Safe Drinking Water Act	Federal Primary Drinking Water Standards – MCLs and MCLGs	42 USC 300f; 40 CFR 141, Subpart O	Establishes drinking water standards for public water systems to protect human health. Includes standards for the following Site contaminants of potential concern: arsenic, benzene, and benzo(a) pyrene. The National Contingency Plan states that MCLs, not MCLGs, are ARARs for usable aquifers.	ARARs for groundwater that could potentially be used for drinking water, where the water will be provided directly to 25 or more people or will be supplied to 15 or more service connections.
Safe Drinking Water Act	Federal Secondary Drinking Water Standards – Secondary MCLs	42 USC 300f; 40 CFR 143	Establishes drinking water standards for public water systems to achieve the aesthetic qualities of drinking water (secondary MCLs).	TBC for groundwater that could potentially be a drinking water source (i.e., achieved as practicable).
Clean Water Act	Federal Ambient Water Quality Criteria	33 USC 1311– 1317; 40 CFR 131	Under Clean Water Act, Section 304(a), minimum criteria are developed for water quality programs established by states. Two kinds of water quality criteria are developed: one for protection of human health, and one for protection of aquatic life. The federal recommended water quality criteria are published on EPA's website: http://water.epa.gov/scitech/swguidance/standards/current/index.cfm	ARARs for surface water if more stringent than promulgated state criteria.
Surface Water Quality Standards	State Ambient Water Quality Criteria	Chapter 90.48 RCW; Chapter 173-201A WAC	Establishes water quality standards for protection of human health and for protection of aquatic life (for both acute and chronic exposure durations).	ARARs for surface water where Washington State has adopted, and EPA has approved, water quality standards.
Model Toxics Control Act	State Soil, Air, Groundwater, and Surface Water Cleanup Standards	Chapter 70.105D RCW; Chapter 173-340 WAC	Establishes cleanup levels for Site groundwater, surface water, soil, and air, including rules for evaluating cross-media protectiveness. MTCA cleanup levels cannot be set at concentrations below natural background.	Promulgated numeric cleanup levels are ARARs for soil, air, groundwater, and surface water. Equations to develop cleanup levels are not ARARs.

## Table 6-1

## Table 6-1 – Potential ARARs, Contaminant-Specific

Bremerton Gas Works Site Bremerton, Washington

Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness
Sediment Management Standards	State Sediment Quality Criteria	Chapters 90.48 & 70.105D RCW; Chapter 173-204 WAC	Establishes both numerical and biological wasting-based standards for the protection of benthic invertebrates in marine sediments. The current rule also defines methods for establishing cleanup levels protective of human health, including protection from risks associated with seafood consumption, analytical considerations, and natural and regional background contamination levels.	SMS cleanup levels will serve as ARARs for the development of sediment cleanup levels.

#### Notes:

ARAR = applicable or relevant and appropriate requirement

CFR = Code of Federal Regulations

EPA = U.S. Environmental Protection Agency

MCL = maximum contaminant level

MCLG = maximum contaminant level goal

MTCA = Model Toxics Control Act

RCW = Revised Code of Washington

SMS = Sediment Management Standards

TBC = to be considered

USC = United States Code

WAC = Washington Administrative Code

## Table 6-2 – Potential ARARs, Location-Specific

**Bremerton Gas Works** 

Bremerton, Washington

Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness
Endangered Species Act	Effects on Endangered Species	16 USC 1531 et seq.; 50 CFR 17	Actions authorized, funded, or carried out by federal agencies may not jeopardize the continued existence of endangered or threatened species or adversely modify or destroy their critical habitats, or must take appropriate mitigation steps.	ARAR for remedial actions that may adversely affect endangered or threatened species or critical habitat present at the Site.
Safe Drinking Water Act	Underground Injection Control, Sole Source Aquifer Program, and Wellhead Protection Program	42 USC 300h–300h- 8; 40 CFR 300.400(g)(4); Chapter 173-160 WAC; WAC 246- 290-135	Resource planning programs designed to prevent contamination of underground sources of drinking water.	The requirements of the City's wellhead protection program are TBCs as a performance standard for groundwater that is a potential drinking water source (i.e., achieved as practicable). (Note that there are no water supply wells near the Site that are currently regulated by the City's program.)
Magnuson- Stevens Fishery Conservation and Management Act	Habitat Impacts	16 USC 1855(b); 50 CFR 600.920	Requires evaluation of impacts on EFH if activities may adversely affect EFH.	ARAR if the remedial action may adversely affect EFH.
Executive Order for Wetlands Protection	Wetlands Impacts	Executive Order 11990 (1977), 40 CFR 6.302(a); 40 CFR 6, App. A	Requires measures to avoid adversely affecting wetlands whenever possible, to minimize wetland destruction, and to preserve the value of wetlands.	ARAR for assessing impacts on wetlands, if any, from the remedial action and for developing appropriate compensatory mitigation.

Notes:

ARAR = applicable or relevant and appropriate requirement

CFR = Code of Federal Regulations

City = City of Bremerton

EFH = essential fish habitat

TBC = to be considered

USC = United States Code

WAC = Washington Administrative Code

Table 6-2

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Table 6-3 – Potential ARARs, Action-Specific

Remedial Activity	Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness
	Solid Waste Disposal Act	Management and Disposal of Solid Waste	42 USC 6901–6917; 40 CFR 257–258	Establishes requirements for the management and disposal of solid wastes.	ARAR for remedial actions that result in upland disposal of excavated or dredged material.
Soil Excavation and Upland Filling	Resource Conservation and Recovery Act (RCRA); Washington Hazardous Waste Management Act and Dangerous Waste Regulations	Generation and Management (Transportation, Treatment, Storage, and Disposal) of Hazardous Waste; Off-Site Land Disposal Considerations	42 USC 6921–22; 40 CFR 260, 261, and 268; Chapter 70.105 RCW; Chapter 173-303 WAC (Chapter 173-307 WAC Pollution Prevention Plans is a TBC)	Defines solid wastes subject to regulation as hazardous wastes.  Requires management of hazardous waste from "cradle to grave" unless exemption applies.  MGP wastes are subject to certain exemptions (e.g, Bevill Amendment provisions)	ARAR for wastes and soils sediments excavated from the Site for off-site disposal, and a TBC for on-site stabilization or containment actions.
	Hazardous Materials Transportation Act	Transport of Hazardous Materials	49 USC 5101 et seq.; 49 CFR 171–177	Establishes requirements for transport of hazardous materials.	ARAR for those hazardous materials (e.g., DNAPL) transported off site.
	Washington Hydraulics Code	Filling of Wetlands	Chapters 75.20 and 77.55 RCW; Chapter 220-110 WAC	Establishes requirements for performing work that would alter existing jurisdictional wetlands.	ARAR if remedial actions such as excavation or capping affect existing jurisdictional wetlands. Remedial actions must result in no net loss of aquatic habitat and function after sequential consideration of avoidance and mitigation, allowing for site-specific evaluations of existing wetland functions.

Table 6-3

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Table 6-3 – Potential ARARs, Action-Specific

Remedial Activity	Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness
Soil Excavation and Upland Filling (Continued)	City of Bremerton Shoreline Master Program and Critical Areas Regulations	Shoreline of Statewide Significance; Fish and Wildlife Habitat Conservation Areas	Chapter 90.58 RCW; Chapter 173-14 WAC; City of Bremerton Ordinance #5299 (effective December 4, 2013); Critical Area Regulations (BMC 20.14) are incorporated into the SMP by reference	Establishes replacement requirements for FWHCAs affected by remedial actions to ensure no net loss of existing ecological function; also establishes requirements for buffers and setbacks from shorelines.	ARAR if remedial actions such as excavation or capping result in impacts within 200 feet of ordinary high water mark or designated FWHCAs. Remedial actions must result in no net loss of aquatic habitat and function after sequential consideration of avoidance and mitigation, allowing for site-specific evaluations of existing shoreline habitat and FWHCAs. Washington's vested rights rule governs which SMP requirements apply in a given circumstance. Substantive requirements of the SMP that were in effect when redevelopment project applications were filed may be ARARs for future redevelopment actions at the Site.
Dredging, Capping, and/or Discharge to Puget Sound	Clean Water Act	Federal Ambient Water Quality Criteria	33 USC 1311–1317; 40 CFR 131	Regulates activities that may result in discharges into navigable waters.	ARAR for control of short-term impacts on surface water due to implementation of remedial actions that include dredging, capping, and discharge of treated water into Puget Sound. Incorporates the substantive provisions of relevant and appropriate Joint Aquatic Resources Permit Application (JARPA), Nationwide Permit, and stormwater regulation requirements.

## Table 6-3

Table 6-3 – Potential ARARs, Action-Specific

Remedial Activity	Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness	
	Surface Water Quality Standards	State Ambient Water Quality Criteria	Chapter 90.48 RCW; Chapter 173-201A WAC	Regulates activities that may result in discharges into navigable waters.	ARAR for control of short-term impacts on surface water sue to implementation of remedial actions that include dredging, capping, and discharge of treated water into Puget Sound. Incorporates the substantive provisions of relevant and appropriate requirements, where Washington State has adopted, and EPA has approved, water quality standards.	
Dredging, Capping, and/or Discharge to Puget Sound	Clean Water Act	Discharge of Materials into Puget Sound	33 USC 1344; 40 CFR 230	Regulates discharge of dredged and fill material into navigable waters of the United States.	ARAR for dredging and capping activities in Puget Sound.	
(Continued)	Fish and Wildlife Coordination Act	Discharge of Materials, Impoundment or Diversion of Waters in Puget Sound  16 USC 662 and 663; 40 CFR 6.302(g)		Requires federal agencies to consider effects on fish and wildlife from projects that may alter a body of water and mitigate or compensate for project-related losses, which include discharges of pollutants to water bodies.	ARAR for in-water remedial actions or if treated water is discharged into Puget Sound.	
	River and Harbors Act  Placement of Structures in Puget Sound		33 USC 401 et seq.; 33 CFR 320–330	Prohibits the unauthorized obstruction or alteration of any navigable water. Establishes requirements for structures or work in, above, or under navigable waters.	ARAR for remedial actions in Puget Sound.	

## Table 6-3

Table 6-3 – Potential ARARs, Action-Specific

Remedial Activity	Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness
Dredging, Capping, and/or Discharge to Puget Sound (Continued)	Washington Hydraulics Code	Filling in Puget Sound	Chapter 75.20 and 77.55 RCW; Chapter 220-110 WAC	Establishes requirements for performing work that would use, divert, obstruct, or change the natural flow or bed of Puget Sound.	ARAR for shoreline excavation, dredging, and/or capping actions. Remedial actions must result in no net loss of aquatic habitat or function after sequential consideration of avoidance and mitigation.
	Federal Clean Air Act; Washington Clean Air Act; Puget Sound Air Clean Air Agency Regulations	Air Emission Discharges	42 USC 7401 et seq.; Chapter 70.94 RCW; Chapter 173-400 WAC; PSCAA Regulation III	Regulates air emission discharges.	ARAR for remedial activities that generate fugitive dust or other air emissions, including treatment operations.
Other Remedial Activities	Historic Preservation Act; Washington Historical Activities Act	Alteration of Historic Properties	16 USC 470 et seq.; 36 CFR 800; Chapter 27 RCW	Requires the identification of historic properties potentially affected by remedial actions, and ways to avoid, minimize, or mitigate such effects. Historic property is any district, site, building, structure, or object included in or eligible for the National Register of Historic Places, including artifacts, records, and material remains related to such a property.	ARAR if historic properties are affected by remedial activities. No historic properties have been identified at the Site to date but could potentially be identified during remedial design.
	Archeological and Historic Preservation Act  Alteration of Historic and Archaeological Properties		16 USC 469a-1	Provides for the preservation of historical and archeological data that may be irreparably lost as a result of a federally approved project and mandates only preservation of the data.	ARAR if historical and archeological resources may be irreparably lost by implementation of remedial activities.

## Table 6-3

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## Table 6-3 – Potential ARARs, Action-Specific

**Bremerton Gas Works** 

Bremerton, Washington

Remedial Activity	Act/Authority	Criteria/Issue	Citation	Brief Description	Applicability/Appropriateness
Other Remedial Activities (Continued)	Native American Graves Protection and Reparation Act	Alteration of American Graves	25 USC 3001–3013; 43 CFR 10	Requires federal agencies and museums that have possession of or control over Native American cultural items (including human remains, associated and unassociated funerary items, sacred objects, and objects of cultural patrimony) to compile an inventory of such items. Prescribes when such federal agencies and museums must return Native American cultural items. "Museums" are defined as any institution or state or local government agency that receives federal funds and has possession of, or control over, Native American cultural items.	ARAR if Native American cultural items are present in an excavation or dredging area.

Notes:

ARAR = applicable or relevant and appropriate requirements

BMC = Bremerton Municipal Code

DNAPL = dense non-aqueous phase liquid

EPA = U.S. Environmental Protection Agency

FWHCA = Fish and Wildlife Habitat Conservation Area

MGP = manufactured gas plant

PSCCA = Puget Sound Clean Air Agency

RCW = Revised Code of Washington

SMP = Shoreline Master Program

TBC = to be considered

USC = United States Code

WAC = Washington Administrative Code

Table 6-3

		EPA Region 5 RCRA Soil Ecological Screening Levels	EPA Ecological Soil Screening Levels - Birds	EPA Ecological Soil Screening Levels - Invertebrates	EPA Ecological Soil Screening Levels - Mammals	EPA Ecological Soil Screening Levels - Plants	EPA Regional Screening Levels (RSLs) - Residential Soil	EPA Regional Screening Levels (RSLs) - Industrial Soil	Scr	s Used for Data eening
Analyte	CAS Number	EPA, 2003	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2013	EPA, 2013	Surface Soil (0-10 feet)	Subsurface Soil (>10 feet)
Alkane Isomers (ug/kg)	CAS IVAIIIDEI	El A, 2003	LI A, 2010	LI A, 2010	LI A, 2010	LI A, 2010	LI A, 2013	LI A, 2013	(0 10 100)	(>10 (cct)
n-Hexane (C6)	110-54-3	IHH	<del>=</del>	<b>₩</b> ₩	<del></del>	<b>E</b>	570000	2600000	570000	570000
Conventionals (mg/kg)	300 00000									
Cyanide, WAD	57-12-5		122	400	nes.		22	140	22	22
Cyanide, total	57-12-5	1.33					·		1.33	
Sulfide	18496-25-8	0.00358	: <del></del> -	4-			:		0.00358	
Metals (mg/kg)										
Antimony	7440-36-0	0.142	a <del></del>	78	0.27		31	410	0.27	31
Arsenic	7440-38-2	5.7	43	-	46	18	0.61	2.4	0.61	0.61
Beryllium	7440-41-7	1.06	-	40	21		160	2000	21	160
Cadmium	7440-43-9	0.00222	0.77	140	0.36	32	70	800	0.36	70
Chromium	7440-47-3	0.4	26	11	34	1220	1		26	
Chromium III	16065-83-1		26	( <del>4.5.</del> )	34		120000	1500000	26	120000
Chromium VI	18540-29-9	II		( <b></b>	130	×==1	0.29	5.6	0.29	0.29
Cobalt	7440-48-4	0.14	120	-	230	13	23	300	13	23
Copper	7440-50-8	5.4	28	80	49	70	3100	41000	28	3100
Lead	7439-92-1	0.0537	11	1700	56	120	400	800	11	400
Manganese	7439-96-5	HB.	4300	450	4000	220	1800	23000	220	1800
Mercury	7439-97-6	0.1					10	43	10	10
Nickel	7440-02-0	13.6	210	280	130	38	1500	20000	38	1500
Selenium	7782-49-2	0.0276	1.2	4.1	0.63	0.52	390	5100	0.52	390
Silver	7440-22-4	4.04	4.2		14	560	390	5100	4.2	390
Thallium	7440-28-0	0.0569	( <del>ii.e</del>	**	<del>är</del> .	<b>Æ</b> .	0.78	10	0.78	0.78
Zinc	7440-66-6	6.62	46	120	79	160	23000	310000	46	23000
Metals, Organic (ug/kg)										
Tributyltin	688-73-3	IMM	(##	HH;		EH.	18000	180000	18000	18000
Polycyclic Aromatic Hydrocarbons (PAHs) (ug/kg)										
1-Methylnaphthalene	90-12-0	122			u=		16000	53000	16000	16000
2-Methylnaphthalene	91-57-6	3240		. <del></del>	<del></del>		230000	2200000	230000	230000
Acenaphthene	83-32-9	682000	×==	-			3400000	33000000	3400000	3400000
Acenaphthylene	208-96-8	682000	:##			##:	3 <del>##</del>	**	682000	
Anthracene	120-12-7	1480000		: <del></del>	i <del>par</del> n	<del></del>	17000000	170000000	17000000	17000000
Benzo(a)anthracene	56-55-3	5210	74E	- (a/a/)	i <del>ww</del> i		150	2100	150	150
Benzo(a) fluoranthene	203-33-8	PE		HH)		44	122	44		
Benzo(a)pyrene	50-32-8	1520					15	210	15	15
Benzo(b)fluoranthene	205-99-2	59800	120	- False V	la inc	wa:	150	2100	150	150
Benzo(b,j)fluoranthene	==.			(All and a second secon			100			
Benzo(b,j,k)fluoranthenes					X		1			

		EPA Region 5 RCRA Soil Ecological Screening Levels	EPA Ecological Soil Screening Levels - Birds	EPA Ecological Soil Screening Levels - Invertebrates	EPA Ecological Soil Screening Levels - Mammals	EPA Ecological Soil Screening Levels - Plants	EPA Regional Screening Levels (RSLs) - Residential Soil	EPA Regional Screening Levels (RSLs) - Industrial Soil		s Used for Data eening
									Surface Soil	Subsurface Soil
Analyte	CAS Number	EPA, 2003	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2013	EPA, 2013	(0-10 feet)	(>10 feet)
Benzo(b,k)fluoranthene	202.42.2	7 <u></u>	> made from		i Ball and i				<del>'H</del> +:	<del>(111</del> )
Benzo(g,h,i)fluoranthene	203-12-3			**	**	H	144	22		
Benzo(g,h,i)perylene	191-24-2	119000	<del></del>	· <del></del> -		;==:		4200		
Benzo(j)fluoranthene	205-82-3	188		1 <u>86</u> 9			380	1300	380	380
Benzo(j,k)fluoranthene	207.00.0	140000					1500		4500	4500
Benzo(k)fluoranthene	207-08-9	148000					1500	21000	1500	1500
Benzofluoranthene (unspecified)	56832-73-6	4720	***	***	H-1	<del></del>	15000	24,0000	45,000	45000
Chrysene	218-01-9	4730	: <del></del>		: <del></del> :	: <del></del> -	15000	210000	15000	15000
Dibenzo(a,e)pyrene	192-65-4	10400			20	/W=1	38	130	38	38
Dibenzo(a,h)anthracene	53-70-3	18400	( <del>****</del>	÷*	<del>""</del>	H-1	15	210	15	15
Fluoranthene	206-44-0 86-73-7	122000 122000	\ <del></del>	·			2300000	22000000 22000000	2300000	2300000
Fluorene	193-39-5	109000					2300000 150	2100	2300000	2300000
Indeno(1,2,3-c,d)pyrene Total HPAH			·=-	18000	1100				150	150
Total LPAH		II <del></del>					1		1100	
				29000	100000	į.			29000	
Naphthalene	91-20-3	99.4	(A)		****		3600	18000	3600	3600
Phenanthrene	85-01-8	45700					4700000	4700000	45700	470000
Pyrene (L.:1)	129-00-0	78500		**************************************	-	- 22	1700000	17000000	1700000	1700000
Total Benzofluoranthenes (b,j,k)		9 <del>=3</del>					·			
Total HPAH		×==	, <del></del>	18000	1100	·=-	×		1100	
Total LPAH				29000	100000	( <del></del>	; <del></del>		29000	
Total PAH		:==	» <del></del>	: •••	: <del>, =</del> :	See S	:	A	7 <b>22</b> 0	
Polychlorinated Biphenyls (PCBs) (ug/kg)  Aroclor 1016	12674 11 2	7779			-	200	2000	21000	2000	2000
and the contraction of the contr	12674-11-2	122	(Callery)	H=1		==	3900	21000	3900	3900
Aroclor 1221	11104-28-2				: 22)		140	540	140	140
Aroclor 1232 Aroclor 1242	11141-16-5 53469-21-9	122	are .	20			140 220	540 740	140	140
Aroclor 1242 Aroclor 1248	12672-29-6		· <del></del>				220	740	220	220
Aroclor 1248 Aroclor 1254	11097-69-1	II				X==1	220	740	220	220 220
Aroclor 1254 Aroclor 1260	The same of the sa			W-3	W.C.	==	220	740	220	
Aroclor 1260 Aroclor 1262	11096-82-5 37324-23-5	100	: <del></del>						220	220
				W21					==	
Aroclor 1268 Total PCB Aroclors	11100-14-4	 0.332	( <del>2)</del>	H=1	<u> </u>	22	220	 740	220	220
Semivolatile Organic Componds (SVOCs) (ug/kg)		0.532	.==		; <del>(22)</del> )	) <del>22</del> 1	220	740	220	220
1,2,4,5-Tetrachlorobenzene	95-94-3	2020					18000	180000	10000	19000
1,2,4-Trichlorobenzene	120-82-1		·	I—==		/	22000	99000	18000	18000
1,2-Dichlorobenzene		11100		(G2)	( == 5)	/ <del></del>			22000	22000
The state of the s	95-50-1	2960	<del></del>	<b>**</b>	WF.	==	1900000	9800000	1900000	1900000
1,3-Dichlorobenzene	541-73-1	37700	: <b></b>			:==	11		37700	

		EPA Region 5 RCRA Soil Ecological Screening Levels	EPA Ecological Soil Screening Levels - Birds	EPA Ecological Soil Screening Levels - Invertebrates	EPA Ecological Soil Screening Levels - Mammals	EPA Ecological Soil Screening Levels - Plants	EPA Regional Screening Levels (RSLs) - Residential Soil	EPA Regional Screening Levels (RSLs) - Industrial Soil		s Used for Data reening
									Surface Soil	Subsurface Soil
Analyte	CAS Number	EPA, 2003	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2013	EPA, 2013	(0-10 feet)	(>10 feet)
1,4-Dichlorobenzene	106-46-7	546				7 <b>4-</b> 0	2400	12000	2400	2400
2,2'-Oxybis (1-chloropropane)	108-60-1	19900	8-	I <del>MH</del> )		-	4600	22000	4600	4600
2,3,4,6-Tetrachlorophenol	58-90-2	199				NN	1800000	18000000	1800000	1800000
2,4,5-Trichlorophenol	95-95-4	14100		(MC)	<u> </u>	( <u>20</u> )	6100000	62000000	6100000	6100000
2,4,6-Trichlorophenol	88-06-2	9940					44000	160000	44000	44000
2,4-Dichlorophenol	120-83-2	87500					180000	1800000	180000	180000
2,4-Dimethylphenol	105-67-9	10	<del>2</del>		<del></del>	<del>(20</del> )	1200000	12000000	1200000	1200000
2,4-Dinitrophenol	51-28-5	60.9		· Amesia	==	S <del>a,</del> — V	120000	1200000	120000	120000
2,4-Dinitrotoluene	121-14-2	1280				200	1600	5500	1600	1600
2,6-Dinitrotoluene	606-20-2	32.8		1		<b>4</b>	330	1200	330	330
2-Chloronaphthalene	91-58-7	12.2		-			6300000	82000000	6300000	6300000
2-Chlorophenol	95-57-8	243				1221	390000	5100000	390000	390000
2-Methylphenol (o-Cresol)	95-48-7	40400	<del>a.</del>	H <del>all</del> ik	( <del>) ()</del>	. <del></del> :	3100000	31000000	3100000	3100000
2-Nitroaniline	88-74-4	74100					610000	6000000	610000	610000
2-Nitrophenol	88-75-5	1600		<b>#</b> #3	-	-			1600	
3,3'-Dichlorobenzidine	91-94-1	646				: <del></del> :	1100	3800	1100	1100
3-Methylphenol & 4-Methylphenol (m&p-Cresol)	1319-77-3				122		6100000	62000000	6100000	6100000
3-Methylphenol (m-Cresol)	108-39-4	3490	÷-	***		-	3100000	31000000	3100000	3100000
3-Nitroaniline	99-09-2	3160				x <b></b> x			3160	200
4-Bromophenyl-phenyl ether	101-55-3			1900		1220				4-
4-Chloro-3-methylphenol	59-50-7	7950				2 <b></b> -	6100000	62000000	6100000	6100000
4-Chloroaniline	106-47-8	1100					2400	8600	2400	2400
4-Methylphenol (p-Cresol)	106-44-5	163000		·	SAR!	(AA)	6100000	62000000	6100000	6100000
4-Nitroaniline	100-01-6	21900		· <del></del> -			24000	86000	24000	24000
4-Nitrophenol	100-02-7	5120			221	2 <b>44</b> 1	:1 <u>==</u>		5120	<del></del>
Acetophenone	98-86-2	300000		HH)	Table 1	<del>=</del>	7800000	100000000	7800000	7800000
Aniline	62-53-3	56.8					85000	300000	85000	85000
Atrazine	1912-24-9		:=-			3 <u></u> 0	2100	7500	2100	2100
Benzaldehyde	100-52-7						7800000	100000000	7800000	7800000
Benzidine	92-87-5					>==>	0.5	7.5	0.5	0.5
Benzo(b)pyridine	91-22-5			¥=1			160	570	160	160
Benzoic acid	65-85-0	1				(55)	240000000	2500000000	240000000	240000000
Benzyl alcohol	100-51-6	65800			220	(HH)	6100000	62000000	6100000	6100000
Biphenyl (1,1'-Biphenyl)	92-52-4		w-	- <del></del>		*	51000	210000	51000	51000
bis(2-Chloroethoxy)methane	111-91-1	302	P6000	9.00-W	1 900000	2000 to	180000	1800000	180000	180000
bis(2-Chloroethyl)ether	111-44-4	23700					210	1000	210	210
bis(2-Ethylhexyl)phthalate	117-81-7	925				/==/	35000	120000	35000	35000
Butylbenzyl phthalate	85-68-7	239					260000	910000	260000	260000

		EPA Region 5 RCRA Soil Ecological Screening Levels	EPA Ecological Soil Screening Levels - Birds	EPA Ecological Soil Screening Levels - Invertebrates	EPA Ecological Soil Screening Levels - Mammals	EPA Ecological Soil Screening Levels - Plants	EPA Regional Screening Levels (RSLs) - Residential Soil	EPA Regional Screening Levels (RSLs) - Industrial Soil	Initial PRGs Used for Data Screening	
									Surface Soil	Subsurface Soil
Analyte	CAS Number	EPA, 2003	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2013	EPA, 2013	(0-10 feet)	(>10 feet)
Caprolactam	105-60-2						3000000	30000000	30000000	3000000
Dibenzofuran	132-64-9		<del></del>		<del>==</del>	**	78000	1000000	78000	78000
Diethyl phthalate	84-66-2	24800				1	49000000	490000000	49000000	49000000
Dimethyl phthalate	131-11-3	734000							734000	
Di-n-butyl phthalate	84-74-2	150				:	6100000	62000000	6100000	6100000
Dinitro-o-cresol (4,6-Dinitro-2-methylphenol)	534-52-1	144					4900	49000	4900	4900
Di-n-octyl phthalate	117-84-0	709000	<del>-</del> -	**	##:	76	610000	6200000	610000	610000
Hexachlorobenzene	118-74-1	199	:	· ( <del>= -</del> ):		<del></del> /	300	1100	300	300
Hexachlorocyclopentadiene	77-47-4	755	:=-				370000	3700000	370000	370000
Hexachloroethane	67-72-1	596	(##	H+:		H#1	12000	43000	12000	12000
Isophorone	78-59-1	139000					510000	1800000	510000	510000
Nitrobenzene	98-95-3	1310	· <del></del>			<del></del>	4800	24000	4800	4800
n-Nitrosodimethylamine	62-75-9	0.0321	/ <del>==</del>	( <del>FE</del> )	A-2		2.3	34	2.3	2.3
n-Nitrosodi-n-propylamine	621-64-7	544		( <b></b> )			69	250	69	69
n-Nitrosodiphenylamine	86-30-6	545		<b>**</b>	H-1	<b>H</b>	99000	350000	99000	99000
Pentachlorophenol	87-86-5	119	2100	31000	2800	5000	890	2700	890	890
Phenol	108-95-2	120000			( <u></u> )	Park send (c)	18000000	180000000	18000000	18000000
Volatile Organic Compounds (VOCs) (ug/kg)						-				
1,1,1,2-Tetrachloroethane	630-20-6	225000			11		1900	9300	1900	1900
1,1,1-Trichloroethane	71-55-6	29800		1227	eres	- Later	8700000	38000000	8700000	8700000
1,1,2,2-Tetrachloroethane	79-34-5	127		(IEEE)			560	2800	560	560
1,1,2-Trichloroethane	79-00-5	28600					1100	5300	1100	1100
1,1,2-Trichlorotrifluoroethane (Freon 113)	76-13-1	. <del>HE</del>	-	**	<del></del> -		43000000	180000000	43000000	43000000
1,1-Dichloroethane	75-34-3	20100	:==	: <del>             </del>		.mmi	3300	17000	3300	3300
1,1-Dichloroethene	75-35-4	8280	:==				240000	1100000	240000	240000
1,2,3-Trichlorobenzene	87-61-6	I <del>M</del>		HH.		<b>₩</b>	49000	490000	49000	49000
1,2,3-Trichloropropane	96-18-4	3360					5	95	5	5
1,2,4-Trimethylbenzene	95-63-6						62000	260000	62000	62000
1,2-Dibromo-3-chloropropane	96-12-8	35.2	, <del></del>	(FA)			5.4	69	5.4	5.4
1,2-Dichloroethane	107-06-2	21200		( <b>——</b> 6			430	2200	430	430
1,2-Dichloroethene, cis-	156-59-2			¥+0		##	160000	2000000	160000	160000
1,2-Dichloroethene, trans-	156-60-5	784	:	-			150000	690000	150000	150000
1,2-Dichloropropane	78-87-5	32700				1441	940	4700	940	940
1,3,5-Trimethylbenzene (Mesitylene)	108-67-8	##	<del>2-</del>	-	4-	-	780000	10000000	780000	780000
1,3-Dichloropropane	142-28-9	n			1244)		1600000	20000000	1600000	1600000
1,3-Dichloropropene, cis-	10061-01-5	398					1		398	
1,3-Dichloropropene, trans-	10061-02-6	398				:			398	
1,4-Dichloro-2-butene, trans-	110-57-6	1					6.9	35	6.9	6.9

#### Table 6-4 - Development of Initial PRGs for Soil

Bremerton Gas Works Site Bremerton, Washington

		EPA Region 5 RCRA Soil Ecological Screening Levels	EPA Ecological Soil Screening Levels - Birds	EPA Ecological Soil Screening Levels - Invertebrates	EPA Ecological Soil Screening Levels - Mammals	EPA Ecological Soil Screening Levels - Plants	EPA Regional Screening Levels (RSLs) - Residential Soil	EPA Regional Screening Levels (RSLs) - Industrial Soil	Sci	s Used for Data reening
									Surface Soil	Subsurface Soil
Analyte	CAS Number	EPA, 2003	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2013	EPA, 2013	(0-10 feet)	(>10 feet)
1,4-Dioxane	123-91-1	2050				~ <b>_</b> _	4900	17000	4900	4900
2-Butanone (MEK)	78-93-3	89600	(##	<del>                                    </del>	( Marie	<b>**</b>	28000000	200000000	28000000	28000000
2-Hexanone (Methyl butyl ketone)	591-78-6	12600			1==1	( <b></b> )	210000	1400000	210000	210000
4-Chlorotoluene	106-43-4			I==0			1600000	20000000	1600000	1600000
4-Isopropyltoluene (4-Cymene)	99-87-6			-						
Acetone	67-64-1	2500					61000000	630000000	61000000	61000000
Acrolein	107-02-8	5270		HE!	HH.	<del></del>	150	650	150	150
Acrylonitrile	107-13-1	23.9	:=-	:I <del>m.</del> k		( <del>a.e.</del> .)	240	1200	240	240
Benzene	71-43-2	255					1100	5400	255	1100
Bromobenzene	108-86-1	1	-	I <del>NE</del> ()	-	-	300000	1800000	300000	300000
Bromochloromethane	74-97-5	1					160000	680000	160000	160000
Bromodichloromethane	75-27-4	540		1900		1200	270	1400	270	270
Bromoform (Tribromomethane)	75-25-2	15900		ĺ			62000	220000	62000	62000
Bromomethane (Methyl bromide)	74-83-9	235		-			7300	32000	7300	7300
Carbon disulfide	75-15-0	94.1	-	H-I	<del></del>	<del></del>	820000	3700000	820000	820000
Carbon tetrachloride (Tetrachloromethane)	56-23-5	2980	1 <del>4.</del>	· Anna			610	3000	610	610
Chlorobenzene	108-90-7	13100					290000	1400000	290000	290000
Chloroethane	75-00-3			(MH)			15000000	61000000	15000000	15000000
Chloroform	67-66-3	1190				X <b>——</b> X	290	1500	290	290
Chloromethane	74-87-3	10400	:	1220			120000	500000	120000	120000
Cyclohexane	110-82-7						7000000	29000000	7000000	7000000
Dibromochloromethane	124-48-1	2050					680	3300	680	680
Dibromomethane	74-95-3	65000	Œ	H-I	ER		25000	110000	25000	25000
Dichlorodifluoromethane	75-71-8	39500	:			× <b>=</b>	94000	400000	94000	94000
Dichloromethane (Methylene chloride)	75-09-2	4050				7 <u></u>	56000	960000	56000	56000
Ethylbenzene	100-41-4	5160	( <del>25</del>		: <del>au</del> s	<del></del>	5400	27000	5400	5400
Ethylene dibromide (1,2-Dibromoethane)	106-93-4	1230					34	170	34	34
Hexachlorobutadiene (Hexachloro-1,3-butadiene)	87-68-3	39.8				9 <b>22</b> 0	6200	22000	6200	6200
Isopropylbenzene (Cumene)	98-82-8			( <del>CO</del> )			2100000	11000000	2100000	2100000
Methyl acetate	79-20-9	1				.==.	78000000	1000000000	78000000	78000000
Methyl iodide (lodomethane)	74-88-4	1230		**					1230	
Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK))	108-10-1	443000					5300000	53000000	5300000	5300000
Methyl tert-butyl ether (MTBE)	1634-04-4		:			12-1	43000	220000	43000	43000
n-Butylbenzene	104-51-8			1 <del>81</del> )		-	3900000	51000000	3900000	3900000
n-Propylbenzene	103-65-1	D==			( <b>100</b>		3400000	21000000	3400000	3400000
o-Xylene	95-47-6				<u></u>	1881	690000	3000000	690000	690000
sec-Butylbenzene	135-98-8		; <del></del>				7800000	10000000	7800000	7800000
Styrene	100-42-5	4690					6300000	36000000	6300000	6300000

#### **Table 6-4 - Development of Initial PRGs for Soil**

Bremerton Gas Works Site Bremerton, Washington

		EPA Region 5 RCRA	EPA Ecological	EPA Ecological Soil	EPA Ecological Soil	EPA Ecological	EPA Regional	EPA Regional Screening Levels		
		Soil Ecological	Soil Screening	Screening Levels -	Screening Levels -	Soil Screening	_	(RSLs) - Industrial	Initial PRG	s Used for Data
		Screening Levels	Levels - Birds	Invertebrates	Mammals	Levels - Plants	Residential Soil	Soil	Scr	eening
									Surface Soil	Subsurface Soil
Analyte	CAS Number	EPA, 2003	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2010	EPA, 2013	EPA, 2013	(0-10 feet)	(>10 feet)
tert-Butylbenzene	98-06-6		S = =			7861)	7800000	100000000	7800000	7800000
Tetrachloroethene (PCE)	127-18-4	9920		HH)		HH.	22000	110000	22000	22000
Toluene	108-88-3	5450		:==:	Î	.=-:	5000000	45000000	5000000	5000000
Total xylene (reported, not calculated)	1330-20-7	10000	22	/AD	22		630000	2700000	630000	630000
Total Xylene		10000	1					1 <del>11</del> 1.		·
Trichloroethene (TCE)	79-01-6	12400				×==	910	6400	910	910
Trichlorofluoromethane (Fluorotrichloromethane)	75-69-4	16400	( <del>***</del>	(HH)	HH.	<del></del>	790000	3400000	790000	790000
Vinyl acetate	108-05-4	12700	:=-			(mm)	970000	4100000	970000	970000
Vinyl chloride	75-01-4	646	2==			7 <b>==</b> 7.	60	1700	60	60

#### Notes:

Compounds frequently associated with MGP-operations.

-- indicates not available

CAS = Chemical Abstract Services

EPA = U.S. Environmental Protection Agency

HPAH = high molecular weight PAH

LPAH = low molecular weight PAH

kg = kilogram

mg = miligram

MGP = manufactured gas plant

ng = nanogram

PRG = preliminary remediation goal

RCRA = Resource Conservation and Recovery Act

RSL = regional screening level

ug = microgram

WAD = Weak Acid Dissociable Cyanide

#### References:

EPA, 2003. EPA Region 5 Resource Conservation Recovery Act (RCRA) Ecological Screening Levels. August 22, 2003.

EPA, 2010. Ecological Soil Screening Levels. Updated October 20, 2010. Cited: January 15, 2014. Available from: http://www.epa.gov/ecotox/ecossl/

EPA, 2013. EPA Regional Screening Levels. November 2013. Available from: http://www.epa.gov/reg3hwmd/risk/human/rb-concentration\_table/Generic\_Tables/docs/master\_sl\_table\_run\_NOV2013.pdf

# **Table 6-5 - Development of Initial PRGs for Groundwater**Bremerton Gas Works Site Bremerton, Washington

		EPA Regional Screening Levels (RSLs) - MCL	EPA Regional Screening Leve (RSLs) - Tapwate
Analyte	CAS Number	EPA, 2013	EPA, 2013
Alkane Isomers (ug/L)	60 to 12 to 120 to 140	1	/as/puntus
n-Hexane (C6)	110-54-3	**	250
Conventionals (mg/L)  Cyanide, free	57-12-5	0.2	0.0014
Sulfide	18496-25-8		
Aetals (ug/L)	10130 23 0		
Antimony	7440-36-0	6	6
Arsenic	7440-38-2	10	0.045
Beryllium	7440-41-7	4	16
Cadmium	7440-43-9	5	6.9
Chromium	7440-47-3	100	S
Chromium III	16065-83-1	**	16000
Chromium VI	18540-29-9 7440-50-8	1300	0.031 620
Copper Lead	7439-92-1	1500	15
Mercury	7439-92-1	2	0.63
Nickel	7440-02-0		300
Selenium	7782-49-2	50	78
Silver	7440-22-4		71
Thallium	7440-28-0	2	0.16
Zinc	7440-66-6		4700
Лetals, Olrganic (ug/L)			
Tributyltin	688-73-3		2.8
olycyclic Aromatic Hydrocarbons (PAHs)(ug/L)	1 22.42.2	I	
1-Methylnaphthalene	90-12-0	==	0.97
2-Methylnaphthalene Acenaphthene	91-57-6 83-32-9		27 400
Acenaphthylene	208-96-8		400
Anthracene	120-12-7		1300
Benzo(a)anthracene	56-55-3		0.029
Benzo(a)pyrene	50-32-8	0.2	0.0029
Benzo(b)fluoranthene	205-99-2		0.029
Benzo(b,k)fluoranthene	;		
Benzo(g,h,i)perylene	191-24-2		
Benzo(j)fluoranthene	205-82-3		0.056
Benzo(k)fluoranthene	207-08-9		0.29
Chrysene	218-01-9		2.9
Dibenzo(a,h)anthracene	53-70-3	==	0.0029
Fluoranthene	206-44-0		630
Fluorene Indeno(1,2,3-c,d)pyrene	86-73-7 193-39-5		220 0.029
Naphthalene	91-20-3		0.029
Phenanthrene	85-01-8		
Pyrene	129-00-0	<u></u>	87
Total Benzofluoranthenes (b,j,k)			
Total HPAH			
Total LPAH	<del></del>		( <del>==</del>
Total PAH			/ <del></del>
Polychlorinated Biphenyls (PCBs) (ug/L)			
Aroclor 1016	12674-11-2		0.96
Aroclor 1221	11104-28-2		0.004
Aroclor 1232 Aroclor 1242	11141-16-5 53469-21-9		0.004 0.034
Aroclor 1242 Aroclor 1248	12672-29-6		0.034
Aroclor 1254	11097-69-1		0.034
Aroclor 1260	11096-82-5		0.034
Aroclor 1262	37324-23-5		
Aroclor 1268	11100-14-4		
Total PCB Aroclors			0.17
emivolatile Organic Compounds (SVOCs) (ug/L)	<u> </u>		-
1,2,4,5-Tetrachlorobenzene	95-94-3		1.2
1,2,4-Trichlorobenzene	120-82-1	70	0.99
1,2-Dichlorobenzene	95-50-1	600	280
1,3-Dichlorobenzene	541-73-1		
1,4-Dichlorobenzene	106-46-7	75	0.42
2,2'-Oxybis (1-chloropropane)	108-60-1		0.31
2,3,4,6-Tetrachlorophenol 2,4,5-Trichlorophenol	58-90-2 95-95-4	- 1989 Z	170 890

# **Table 6-5 - Development of Initial PRGs for Groundwater**Bremerton Gas Works Site Bremerton, Washington

Analyte	CAS Number	EPA Regional Screening Levels (RSLs) - MCL EPA, 2013	EPA Regional Screening Levels (RSLs) - Tapwater EPA, 2013
2,4,6-Trichlorophenol	88-06-2	LFA, 2013	3.5
2,4-Dichlorophenol	120-83-2		35
2,4-Dimethylphenol	105-67-9		270
2,4-Dinitrophenol	51-28-5		30
2,4-Dinitrotoluene	121-14-2	<del> </del>	0.2
2,6-Dinitrotoluene	606-20-2		0.042
2-Chloronaphthalene	91-58-7		550
2-Chlorophenol	95-57-8		71
2-Methylphenol (o-Cresol)	95-48-7		720
2-Nitroaniline	88-74-4		150
2-Nitrophenol	88-75-5		
3,3'-Dichlorobenzidine  3-Methylphenol & 4-Methylphenol (m&p-Cresol)	91-94-1 1319-77-3		0.11 1400
3-Methylphenol (m-Cresol)	108-39-4		720
3-Nitroaniline	99-09-2		720
4-Bromophenyl-phenyl ether	101-55-3		()——:
4-Chloro-3-methylphenol	59-50-7		1100
4-Chloroaniline	106-47-8		0.32
4-Methylphenol (p-Cresol)	106-44-5		1400
4-Nitroaniline	100-01-6		3.3
4-Nitrophenol	100-02-7		1-2-
Acetophenone	98-86-2		1500
Aniline	62-53-3		12
Atrazine	1912-24-9	3	0.26
Benzaldehyde	100-52-7		1500
Benzidine	92-87-5		0.000092
Benzoic acid	65-85-0		58000
Benzyl alcohol	100-51-6		1500
Biphenyl (1,1'-Biphenyl)	92-52-4		0.83
bis(2-Chloroethoxy)methane	111-91-1	EE	46
bis(2-Chloroethyl)ether	111-44-4		0.012
bis(2-Ethylhexyl)phthalate	117-81-7	6	4.8
Butylbenzyl phthalate  Caprolactam	85-68-7 105-60-2		7700
Dibenzofuran	132-64-9		5.8
Diethyl phthalate	84-66-2		11000
Dimethyl phthalate	131-11-3		
Di-n-butyl phthalate	84-74-2		670
Dinitro-o-cresol (4,6-Dinitro-2-methylphenol)	534-52-1		1.2
Di-n-octyl phthalate	117-84-0		160
Hexachlorobenzene	118-74-1	1	0.042
Hexachlorocyclopentadiene	77-47-4	50	22
Hexachloroethane	67-72-1		0.79
Isophorone	78-59-1		67
Nitrobenzene	98-95-3		0.12
n-Nitrosodimethylamine	62-75-9		0.00042
n-Nitrosodi-n-propylamine	621-64-7		0.0093
n-Nitrosodiphenylamine	86-30-6		10
Pentachlorophenol	87-86-5	1	0.035
Phenol	108-95-2		4500
Volatile Organic Compounds (VOCs) (ug/L) 1,1,1,2-Tetrachloroethane	630-20-6		0.5
1,1,1,2-Tetrachloroethane 1,1,1-Trichloroethane	71-55-6	200	7500
1,1,2,2-Tetrachloroethane	71-55-6	200	0.066
1,1,2-Trichloroethane	79-34-5	5	0.24
1,1,2-Trichloroethane (Freon 113)	76-13-1		53000
1,1-Dichloroethane	75-34-3		2.4
1,1-Dichloroethene	75-35-4	7	260
1,2,3-Trichlorobenzene	87-61-6		5.2
1,2,3-Trichloropropane	96-18-4		0.00065
1,2,4-Trimethylbenzene	95-63-6	120	15
1,2-Dibromo-3-chloropropane	96-12-8	0.2	0.00032
1,2-Dichloroethane	107-06-2	5	0.15
1,2-Dichloroethene, cis-	156-59-2	70	28
1,2-Dichloroethene, trans-	156-60-5	100	86
1,2-Dichloropropane	78-87-5	5	0.38
1,3,5-Trimethylbenzene (Mesitylene)	108-67-8		87
1,3-Dichloropropane	142-28-9		290

Table 6-5

#### **Table 6-5 - Development of Initial PRGs for Groundwater**

Bremerton Gas Works Site Bremerton, Washington

		EPA Regional Screening Levels (RSLs) - MCL	EPA Regional Screening Level (RSLs) - Tapwate
Analyte	CAS Number	EPA, 2013	EPA, 2013
1,3-Dichloropropene, cis-	10061-01-5		
1,3-Dichloropropene, trans-	10061-02-6		
1,4-Dichloro-2-butene, trans-	110-57-6		0.0012
1,4-Dioxane	123-91-1		0.67
2-Butanone (MEK)	78-93-3		4900
2-Chlorotoluene	95-49-8		180
2-Hexanone (Methyl butyl ketone)	591-78-6		34
4-Chlorotoluene	106-43-4		190
4-Isopropyltoluene (4-Cymene)	99-87-6		
Acetone	67-64-1		12000
Acrolein	107-02-8		0.041
Acrylonitrile	107-13-1		0.045
Benzene	71-43-2	5	0.39
Bromobenzene	108-86-1		54
Bromochloromethane	74-97-5		83
Bromodichloromethane	75-27-4	80	0.12
Bromoform (Tribromomethane)	75-25-2	80	7.9
Bromomethane (Methyl bromide)	74-83-9		7
Carbon disulfide	75-15-0		720
Carbon tetrachloride (Tetrachloromethane)	56-23-5	5	0.39
Chlorobenzene	108-90-7	100	72
Chloroethane	75-00-3		21000
Chloroform	67-66-3	80	0.19
Chloromethane	74-87-3		190
Cyclohexane	110-82-7		13000
Dibromochloromethane	124-48-1	80	0.15
Dibromomethane	74-95-3		7.9
Dichlorodifluoromethane	75-71-8		190
Dichloromethane (Methylene chloride)	75-09-2	5	9.9
Ethylbenzene	100-41-4	700	1.3
Ethylene dibromide (1,2-Dibromoethane)	106-93-4	0.05	0.0065
Hexachlorobutadiene (Hexachloro-1,3-butadiene)	87-68-3		0.26
Isopropylbenzene (Cumene)	98-82-8		390
Methyl acetate	79-20-9		16000
Methyl iodide (lodomethane)	74-88-4		
Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK))	108-10-1		1000
Methyl tert-butyl ether (MTBE)	1634-04-4		12
n-Butylbenzene	104-51-8		780
n-Propylbenzene	103-65-1		530
o-Xylene	95-47-6		190
sec-Butylbenzene	135-98-8		1600
Styrene	100-42-5	100	1100
tert-Butylbenzene	98-06-6		510
Tetrachloroethene (PCE)	127-18-4	5	9.7
Toluene	108-88-3	1000	860
Total xylene (reported, not calculated)	1330-20-7	10000	190
Total Xylene	1550-20-7		
Trichloroethene (TCE)	79-01-6	5	0.44
Trichlorofluoromethane (Fluorotrichloromethane)	75-69-4		1100
Vinyl acetate	108-05-4		410
Vinyl acetate Vinyl chloride	75-01-4	2	0.015

#### Notes:

Compounds frequently associated with MGP-operations.

'-- indicates not available

CAS = Chemical Abstract Services

EPA = U. S. Environmental Protection Agency

L = liter

MCL = maximum contaminant level

mg = miligram

MGP = manufactured gas plant

ng = nanogram

 ${\sf PRG} = {\sf preliminary} \ {\sf remediation} \ {\sf goal}$ 

RSL = regional screening level

ug = microgram

#### References:

EPA, 2013. EPA Regional Screening Levels. November 2013. Available from: http://www.epa.gov/reg3hwmd/risk/human/rb-concentration\_table/Generic\_Tables/docs/master\_sl\_table\_run\_NOV2013.pdf

## **Table 6-6 - Development of Initial PRGs for Sediment**Bremerton Gas Works Site Bremerton, Washington

Analyte	CAS Number	SMS Marine Sediment Cleanup Objective (SCO <sup>1</sup> /LAET <sup>2</sup> ) DOE, 2013	SMS Marine Cleanup Screening Level (CSL <sup>1</sup> /2LAET <sup>2</sup> ) DOE, 2013	EPA Region 5 RCRA Sediment Ecological Screening Levels EPA, 2003	EPA Region 3 BTAG Marine Sediment Screening Benchmarks EPA, 2006	Effects Range-Low (ERL) Long et al., 1995	Effects Range- Median (ERM) Long et al., 1995	Initial PRGs Used for Data Screening
Alkane Isomers (ug/kg)	C/10 (4dillioci		,		2.1, 2.1.1		8,	Screening
n-Hexane (C6)	110-54-3	:			39.60	-24	-	39.6
Conventionals (mg/kg)		ı						
Cyanide, WAD	57-12-5			0.0001	0.1			0.1
Cyanide, total Sulfide	57-12-5 18496-25-8			0.0001	130	),TT		0.0001 130
Metals (mg/kg)	10430 23 8	,			130			150
Antimony	7440-36-0	:==	1440		2	172100		2
Arsenic	7440-38-2	57	93	9.79	7.24	8.2	70	57
Beryllium	7440-41-7							www.
Cadmium	7440-43-9	5.1	6.7	0.99	0.68	1.2	9.6	5.1
Chromium	7440-47-3	260	270	43,4	52.3	81	370	260
Chromium III	16065-83-1	.55	===	185	(AAA)	1,51,57		
Chromium VI Copper	18540-29-9 7440-50-8	 390	390	 31.6	18.7	 34	270	390
Lead	7439-92-1	450	530	35.8	30.2	46.7	218	450
Mercury	7439-97-6	0.41	0.59	0.174	0.13	0.15	0.71	0.41
Nickel	7440-02-0	CHAP CONTRACTOR	(Walk)	22.7	15.9	20.9	51.6	20.9
Selenium	7782-49-2	1555	, <del></del>		2	1555	==1	2
Silver	7440-22-4	6.1	6.1	0.5	0.73	1	3.7	6.1
Thallium	7440-28-0	্যন্ত	( <del>ADDR</del> )		100.000	1505		
Zinc	7440-66-6	410	960	121	124	150	410	410
Metals, Organic (ug/kg) Tributyltin	688-73-3							
Polycyclic Aromatic Hydrocarbons (PAHs) (ug/kg)  1-Methylnaphthalene	90-12-0	20	111	20	<u> </u>	400	22	
2-Methylnaphthalene	91-57-6	670	670	20.2	20.2	 70	670	670
Acenaphthene	83-32-9	500	500	6.71	6.71	16	500	500
Acenaphthylene	208-96-8	1300	1300	5.87	5.87	44	640	1300
Anthracene	120-12-7	960	960	57.2	46.9	85.3	1100	960
Benzo(a)anthracene	56-55-3	1300	1600	108	74.8	261	1600	1300
Benzo(a) pyrene	50-32-8	1600	1600	150	88.8	430	1600	1600
Benzo(b)fluoranthene	205-99-2			10400				10400
Benzo(b,k)fluoranthene		204	(4.5)	MA.	27.2	THE		27.2
Benzo(g,h,i)perylene	191-24-2	670	720	170	170	· <del>···</del>		670
Benzo(j)fluoranthene Benzo(k)fluoranthene	205-82-3		(A15) (A15)	240	240	188		240
Chrysene	218-01-9	1400	2800	166	108	384	2800	1400
Dibenzo(a,h)anthracene	53-70-3	230	230	33	6.22	63.4	260	230
Fluoranthene	206-44-0	1700	2500	423	113	600	5100	1700
Fluorene	86-73-7	540	540	77.4	21.2	19	540	540
Indeno(1,2,3-c,d)pyrene	193-39-5	600	690	200	17	ie#		600
Naphthalene	91-20-3	2100	2100	176	34.6	160	2100	2100
Phenanthrene	85-01-8	1500	1500	204	86.7	240	1500	1500
Pyrene	129-00-0	2600	3300	195	153	665	2600	2600
Total Benzofluoranthenes (b,j,k) Total HPAH	-	3200 12000	3600 17000		 655	1700	9600	3200 12000
Total LPAH		5200	5200	45	312	552	3160	5200
Total PAH	-				2900	4022	44792	4022
Polycyclic Aromatic Hydrocarbons (PAHs) (mg/kg-OC)					7 (4) (4) (4) (4) (4)			
2-Methylnaphthalene	91-57-6	38	64					38
Acenaphthene	83-32-9	16	57	20	W.			16
Acenaphthylene	208-96-8	66	66					66
Anthracene	120-12-7	220	1200	<del>/2</del>	<del>2.</del>	35		220
Benzo(a)anthracene Benzo(a)pyrene	56-55-3 50-32-8	110 99	270 210					110 99
Benzo(g,h,i)perylene	191-24-2	31	78					31
Chrysene	218-01-9	110	460		<del>22</del>			110
Dibenzo(a,h)anthracene	53-70-3	12	33					12
Fluoranthene	206-44-0	160	1200	. <del>25</del>	77. 10.	l ex		160
Fluorene	86-73-7	23	79					23
Indeno(1,2,3-c,d)pyrene	193-39-5	34	88	<del>15</del>	20	<del></del>		34
Naphthalene Phenanthrene	91-20-3 85-01-8	99 100	170 480					99 100
Pyrene Pyrene	129-00-0	1000	1400					1000
Total Benzofluoranthenes (b,j,k)	129-00-0	230	450		**			230
Total HPAH		960	5300			1==		960
Total LPAH		370	780	44	22	**		370
Polychlorinated Biphenyls (PCBs) (ug/kg)								
Aroclor 1016	12674-11-2	)##	(58)	<b></b>	<del>=</del>	=	227	
Aroclor 1221	11104-28-2							
Arcelor 1242	11141-16-5	1000 N	(200)	I 330	=		7007	
Aroclor 1242 Aroclor 1248	53469-21-9 12672-29-6							
Aroclor 1248 Aroclor 1254	11097-69-1				63.3			63.3
Aroclor 1260	11096-82-5	——————————————————————————————————————		<del>-</del>		=	#	
Aroclor 1262	37324-23-5	( <del></del>			E-H	(FF	1	
Aroclor 1268	11100-14-4	-	<del>500</del>	*	<del></del>	#		
Total PCB Aroclors		130	1000	59.8	40	22.7	180	130
Polychlorinated Biphenyls (PCBs) (mg/kg-OC) Total PCB Aroclors		12	65					12
Semivolatile Organic Compounds (SVOCs)(ug/kg)								
1,2,4,5-Tetrachlorobenzene	95-94-3			1252	47000			47000
1,2,4-Trichlorobenzene	120-82-1	31	51	5062	473			31
1,2-Dichlorobenzene	95-50-1	35	50	294	989			35
1,3-Dichlorobenzene 1,4-Dichlorobenzene	541-73-1	110		1315	842	##	**	842
WE THE CHARLES CONTROL OF THE CONTRO	106-46-7	110	110	318	460			110
2,2'-Oxybis (1-chloropropane) 2,3,4,6-Tetrachlorophenol	108-60-1 58-90-2			 129	284			284
2,4,5-Trichlorophenol	95-95-4	=			819			819
2,4,6-Trichlorophenol	88-06-2			208	2650			2650
	The second second second second	lis .		1000 N TK	moves (TSCT)			

## **Table 6-6 - Development of Initial PRGs for Sediment**Bremerton Gas Works Site Bremerton, Washington

Analyte	CAS Number	SMS Marine Sediment Cleanup Objective (SCO <sup>1</sup> /LAET <sup>2</sup> ) DOE, 2013	SMS Marine Cleanup Screening Level (CSL <sup>1</sup> /2LAET <sup>2</sup> ) DOE, 2013	EPA Region 5 RCRA Sediment Ecological Screening Levels EPA, 2003	EPA Region 3 BTAG Marine Sediment Screening Benchmarks EPA, 2006	Effects Range-Low (ERL) Long et al., 1995	Effects Range- Median (ERM) Long et al., 1995	Initial PRGs Used for Data Screening
2,4-Dichlorophenol	120-83-2			81.7	117			117
2,4-Dimethylphenol	105-67-9	29	29	304	29	1 <del>2.7</del>	==:	29
2,4-Dinitrophenol	51-28-5			6.21				6.21
2,4-Dinitrotoluene	121-14-2	.553		14.4	41.6	1.75.77	.=e.i	41.6
2,6-Dinitrotoluene	606-20-2	-	-	39.8	-			39.8
2-Chloronaphthalene	91-58-7		77.	417	(52)	1.515		417
2-Chlorophenol	95-57-8			31.9	344			344
2-Methylphenol (o-Cresol)	95-48-7	63	63	55.4		V <del>. (1</del>	<del></del>	63
2-Nitroaniline	88-74-4 88-75-5							
2-Nitrophenol 3.3'-Dichlorobenzidine	91-94-1			 127	2060	line line		2060
3-Methylphenol & 4-Methylphenol (m&p-Cresol)	1319-77-3	777				357		2060
3-Methylphenol (m-Cresol)	108-39-4			52.4				52.4
3-Nitroaniline	99-09-2							32. <del>1</del>
4-Bromophenyl-phenyl ether	101-55-3			1550	1230			1230
4-Chloro-3-methylphenol	59-50-7	ces.		388	(Res)	1.555		388
4-Chloroaniline	106-47-8		-	146	-			146
4-Methylphenol (p-Cresol)	106-44-5	670	670	20.2	670	5 <del>777</del>	(লক্ড)	670
4-Nitroaniline	100-01-6	S==	544		(500)			
4-Nitrophenol	100-02-7	455	3 <del>777</del> 0	13.3	(West)	i <del>na</del>	. <del>-</del>	13.3
Acetophenone	98-86-2	:==			(##)			
Aniline	62-53-3	- 655	-55	0.31	555	.555		0.31
Atrazine	1912-24-9	-	144		6.62			6.62
Benzaldehyde	100-52-7	una	- <del>1000</del> 0	(BD)	170 MIN	1959	. <del></del>	
Benzidine Penzidi acid	92-87-5	 650						 CEO
Benzoic acid Benzyl alcohol	65-85-0 100-51-6	650 57	650 73	1.04	650			650 57
Biphenyl (1,1'-Biphenyl)	92-52-4		/3 	1.04	1220			1220
bis(2-Chloroethoxy)methane	92-52-4		-55	==				1220
bis(2-Chloroethyl)ether	111-91-1			3520		100		3520
bis(2-Ethylhexyl)phthalate	117-81-7	1300	3100	182	182	1.0000		1300
Butylbenzyl phthalate	85-68-7	63	900	1970	16800			63
Caprolactam	105-60-2				-			
Dibenzofuran	132-64-9	540	540	449	7300	1970	<b>45</b> .	540
Diethyl phthalate	84-66-2	200	1200	295	218	1200		200
Dimethyl phthalate	131-11-3	71	160	i <del>se</del>	(70.00)	150	(FB)	71
Di-n-butyl phthalate	84-74-2	1400	5100	1114	1160			1400
Dinitro-o-cresol (4,6-Dinitro-2-methylphenol)	534-52-1			104	(5.5)	177		104
Di-n-octyl phthalate	117-84-0	6200	6200	40600	100			6200
Hexachlorobenzene	118-74-1	22	70	20	20	li <del>cato</del> r:	.=q;	22
Hexachlorocyclopentadiene	77-47-4			901 584	139 804			139
Hexachloroethane Isophorone	67-72-1 78-59-1	)		432				804 432
Nitrobenzene	98-95-3			145				145
n-Nitrosodimethylamine	62-75-9		Switch Control of the		100 may			
n-Nitrosodi-n-propylamine	621-64-7		-		7.7	1000		
n-Nitrosodiphenylamine	86-30-6	28	40		422000			28
Pentachlorophenol	87-86-5	360	690	23000	7970	1.555		360
Phenol	108-95-2	420	1200	49.1	420	1-4		420
Semivolatile Organic Compounds (SVOCs) (mg/kg-OC)				10				
1,2,4-Trichlorobenzene	120-82-1	0.81	1.8		M-10.			0.81
1,2-Dichlorobenzene	95-50-1	2.3	2.3	. <del> </del>	( <del>la ve</del> s)	LONG.	.=q <sub>0</sub>	2.3
1,4-Dichlorobenzene	106-46-7	3.1	9					3.1
bis(2-Ethylhexyl)phthalate Butylbenzyl phthalate	117-81-7 85-68-7	47 4.9	78 64					47 4.9
Dibenzofuran	132-64-9	15	58		==	VET		15
Diethyl phthalate	84-66-2	61	110					61
Dimethyl phthalate	131-11-3	53	53					53
Di-n-butyl phthalate	84-74-2	220	1700	2000	00 (20)	1.000	50105	220
Di-n-octyl phthalate	117-84-0	58	4500			1		58
Hexachlorobenzene	118-74-1	0.38	2.3	==	gar)			0.38
n-Nitrosodiphenylamine	86-30-6	11	11:	20.	: <del>7.2</del> 0	LETTER.		11
Volatile Organic Compounds (VOCs) (ug/kg)								
1,1,1,2-Tetrachloroethane	630-20-6	-cna	-570	20.	575	ina.		
1,1,1-Trichloroethane	71-55-6			213	856			856
1,1,2,2-Tetrachloroethane	79-34-5			850	202			202
1,1,2-Trichloroethane	79-00-5		-	518	570			570
1,1,2-Trichlorotrifluoroethane (Freon 113) 1,1-Dichloroethane	76-13-1 75-34-3			 0.575	7.2. 			 0 575
1,1-Dichloroethane 1,1-Dichloroethene	75-34-3 75-35-4		-	0.575 19.4	 2780			0.575 2780
1,2,3-Trichlorobenzene	75-35-4 87-61-6			19.4 	858			858
1,2,3-Trichloropropane	96-18-4		_			100		
1,2,4-Trimethylbenzene	95-63-6		M092	1900	1905		514051 	
	96-12-8							
1,2-Dibromo-3-chloropropane	96-12-8		<b></b>	260				260
	107-06-2	See.		200	:==::			
1,2-Dibromo-3-chloropropane						155		
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane	107-06-2		<del>                                     </del>	1				1050
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis-	107-06-2 156-59-2			20		inc.		1050 333
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans-	107-06-2 156-59-2 156-60-5			 654	 1050			
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9			 654 333	 1050 	 		333
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis-	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5	200 200 200 200 200		 654 333  	1050			333  
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans-	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6			 654 333   	1050			333
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans-	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6	100 100 100 100 100 100		 654 333    	1050   	200 200 200 200 200 200 200 200 200 200		333
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans- 1,4-Dioxane	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6 123-91-1	200 200 200 200 200 200 200 200 200 200	100 100 100 100 100 100 100 100 100 100	 654 333      119	 1050    	200 200 200 200 200 200 200 200 200 200		333     119
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans- 1,4-Dioxane 2-Butanone (MEK)	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6 123-91-1 78-93-3		100 100 100 100 100 100 100 100 100 100	 654 333     119 42.4	1050 	200 200 200 200 200 200 200 200 200 200		333    119 42.4
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans- 1,4-Dioxane 2-Butanone (MEK) 2-Chlorotoluene	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6 123-91-1 78-93-3 95-49-8	100 100 100 100 100 100 100 100 100 100		 654 333     119 42.4		200 - 200 -		333     119 42.4
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans- 1,4-Dioxane 2-Butanone (MEK) 2-Chlorotoluene 2-Hexanone (Methyl butyl ketone)	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6 123-91-1 78-93-3 95-49-8 591-78-6			 654 333    119 42.4  58.2	1050 			333    119 42.4  58.2
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans- 1,4-Dioxane 2-Butanone (MEK) 2-Chlorotoluene 2-Hexanone (Methyl butyl ketone) 4-Chlorotoluene	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6 123-91-1 78-93-3 95-49-8 591-78-6 106-43-4							333    119 42.4  58.2
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans- 1,4-Dioxane 2-Butanone (MEK) 2-Chlorotoluene 2-Hexanone (Methyl butyl ketone) 4-Chlorotoluene 4-Isopropyltoluene (4-Cymene)	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6 123-91-1 78-93-3 95-49-8 591-78-6 106-43-4 99-87-6				1050			333    119 42.4  58.2
1,2-Dibromo-3-chloropropane 1,2-Dichloroethane 1,2-Dichloroethene, cis- 1,2-Dichloroethene, trans- 1,2-Dichloropropane 1,3,5-Trimethylbenzene (Mesitylene) 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropene, cis- 1,3-Dichloropropene, trans- 1,4-Dichloro-2-butene, trans- 1,4-Dioxane 2-Butanone (MEK) 2-Chlorotoluene 2-Hexanone (Methyl butyl ketone) 4-Chlorotoluene	107-06-2 156-59-2 156-60-5 78-87-5 108-67-8 142-28-9 10061-01-5 10061-02-6 110-57-6 123-91-1 78-93-3 95-49-8 591-78-6 106-43-4							333    119 42.4  58.2

#### Table 6-6 - Development of Initial PRGs for Sediment

Bremerton Gas Works Site Bremerton, Washington

Analyte	CAS Number	SMS Marine Sediment Cleanup Objective (SCO <sup>1</sup> /LAET <sup>2</sup> ) DOE, 2013	SMS Marine Cleanup Screening Level (CSL <sup>1</sup> /2LAET <sup>2</sup> ) DOE, 2013	EPA Region 5 RCRA Sediment Ecological Screening Levels EPA, 2003	EPA Region 3 BTAG Marine Sediment Screening Benchmarks EPA, 2006	Effects Range-Low (ERL) Long et al., 1995	Effects Range- Median (ERM) Long et al., 1995	Initial PRGs Used for Data Screening
Benzene	71-43-2			142	137			137
Bromobenzene	108-86-1	455			(A-45)			
Bromochloromethane	74-97-5	See.			(mm).			
Bromodichloromethane	75-27-4	155	100		1000	155		
Bromoform (Tribromomethane)	75-25-2	5 <del></del>	-	492	1310			1310
Bromomethane (Methyl bromide)	74-83-9	-10000	- <del>77.77</del> .	1.37	NVIDE:	1571659	57-00E	1.37
Carbon disulfide	75-15-0	5 <del>14</del>	-	23.9	0.851			0.851
Carbon tetrachloride (Tetrachloromethane)	56-23-5	No.		1450	7240	II <del>CC</del>	550	7240
Chlorobenzene	108-90-7			291	162			162
Chloroethane	75-00-3	1 Labore		i assi	1 <del>0.00</del>	1555	.ee/	
Chloroform	67-66-3	: www	1441	121				121
Chloromethane	74-87-3	15000	100000		NAME:	II <del>MM</del> .	.T.	
Cyclohexane	110-82-7	3 <del></del>	144					221
Dibromochloromethane	124-48-1	1 (100)			1777	LEG.		.==1
Dibromomethane	74-95-3	; max						
Dichlorodifluoromethane	75-71-8	· LEGGE			1 <del>555</del> 1	LEG.	EQ.	
Dichloromethane (Methylene chloride)	75-09-2	; max	1441	159				159
Ethylbenzene	100-41-4	VAC	: <del>7.7.7.</del> 1	175	305	v.m.n	<del></del>	305
Ethylene dibromide (1,2-Dibromoethane)	106-93-4	:==	1440					221
Hexachlorobutadiene (Hexachloro-1,3-butadiene)	87-68-3	11	120	26.5	(No. 100)	II <del>AA</del>	550	11
Isopropylbenzene (Cumene)	98-82-8	3 <b></b>			86			86
Methyl acetate	79-20-9	1 Table		I ASSI	1 <del>555</del> 5	155	EQ.	
Methyl iodide (Iodomethane)	74-88-4	:						
Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK))	108-10-1	1500	1970	25.1	No.	UMG.	.TE2	25.1
Methyl tert-butyl ether (MTBE)	1634-04-4		1440					<del></del>
n-Butylbenzene	104-51-8	1 1000			1777	LEG.		. <del></del> l
n-Propylbenzene	103-65-1	( max						
o-Xylene	95-47-6	1 <del></del>			1 <del>555</del> 1	LEG.	==0	
sec-Butylbenzene	135-98-8	; max	1441					
Styrene	100-42-5	-155	-570	254	7070	185	.en:	7070
tert-Butylbenzene	98-06-6	:==	1440					<del></del>
Tetrachloroethene (PCE)	127-18-4	1000		990	190	LEG.		190
Toluene	108-88-3			1220	1090			1090
Total xylene (reported, not calculated)	1330-20-7			433		7.7	77:	433
Total Xylene				433				433
Trichloroethene (TCE)	79-01-6	-LEED	-55.	112	8950	uro.	.==o	8950
Trichlorofluoromethane (Fluorotrichloromethane)	75-69-4	c##						
Vinyl acetate	108-05-4	-155	-55	13	1777	una.	.=a;	13
Vinyl chloride	75-01-4	Sime.	122	202				202
Volatile Organic Compounds (VOCs) (mg/kg-OC)								
Hexachlorobutadiene (Hexachloro-1,3-butadiene)	87-68-3	3.9	6.2					3.9

#### Notes:

Compounds frequently associated with MGP-operations

- '-- indicates not available
- 1 = This criteria will be used when total organic carbon (TOC) is between 0.5% to 5%.
- 2 = This criteria will be used when total organic carbon (TOC) is less than 0.5% or greater than 5%.

2LAET = Second Lowest Apparent Effects Threshold

BTAG = Biological Technical Assistance Group CAS = Chemical Abstract Services

CSL = Cleanup Screening Level

DOE = Washington Department of Ecology

EPA = United States Environmental Protection Agency

kg = kilogram

LAET = Lowest Apparent Effects Threshold

mg = miligram

MGP = Manufactured Gas Plant

ng = nanogram

OC = organic carbon

PRG = preliminary remediation goal

RCRA = Resource Conservation and Recovery Act

SCO = Sediment Cleanup Objective

SMS = Sediment Management Standards

ug = microgram

#### References:

Ecology, 2013. Sediment Management Standards, Chapter 173-204 WAC: Final Rule February 22, 2013. September 1, 2013.

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Analyte	CAS Number	EPA Region 3 BTAG Marine Water Screening Benchmarks EPA, 2006	National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CCC (chronic) <sup>1</sup> EPA, 2013	National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CMC (acute) <sup>1</sup> EPA, 2013	EPA Region 5 RCRA - Ecological Screening Levels - Water EPA, 2003	National Recommended Water Quality Criteria - Human Health for the Consumption of Organisms EPA, 2013	Initial PRGs Used for Data Screening
Alkane Isomers (ug/L)	110 [4.2	0.50	1			1	
n-Hexane (C6) Conventionals (mg/L)	110-54-3	0.58			944	7880	0.58
Cyanide, free Cyanide, total	57-12-5 57-12-5	0.001	0.001	0.001	0.0052	0.14	0.001 0.14
Sulfide	18496-25-8	S <del>=</del>		22			W-1
Metals (ug/L) Antimony	7440-36-0	500	(57)		80	640	640
Arsenic Beryllium	7440-38-2 7440-41-7	12.5 (a) 0.66	36	69 	148 3.6	0.14	0.14 0.66
Cadmium	7440-43-9	0.12 (a)	8.8	40	0.15	(1209)	8.8
Chromium Chromium III	7440-47-3 16065-83-1	57.5 56 (a)	1555 1555		42	100	57.5 56
Chromium VI Copper	18540-29-9 7440-50-8	1.5 (a) 3.1	50 3.1	1100 4.8	1.58	(SEC)	50 3.1
Lead	7439-92-1	8.1	8.1	210	1.17	100	8.1
Mercury Nickel	7439-97-6 7440-02-0	0.016 (a) 8.2	0.94 8.2	1.8 74	0.0013 28.9	4600	0.94 8.2
Selenium Silver	7782-49-2 7440-22-4	71 0.23	71	290 1.9	5 0.12	4200	71 0.23
Thallium	7440-22-4	21.3	1980	MAY.	10	0.47	0.47
Zinc Metals, Organic (ug/L)	7440-66-6	81	81	90	65.7	26000	81
Tributyltin	688-73-3	0.001 (a)	0.0074	0.42		( <del>C</del> )	0.0074
Polycyclic Aromatic Hydrocarbons (PAHs) (ug/L)  1-Methylnaphthalene	90-12-0	2.1			.==		2.1
2-Methylnaphthalene Acenaphthene	91-57-6 83-32-9	4.2 6.6		-	330 38	990	4.2 990
Acenaphthylene	208-96-8				4840		4840
Anthracene Benzo(a)anthracene	120-12-7 56-55-3	0.18 0.018	122		0.035 0.025	40000 0.018	40000 0.018
Benzo(a)pyrene Benzo(b)fluoranthene	50-32-8 205-99-2	0.015			0.014 9.07	0.018 0.018	0.018 0.018
Benzo(b,k)fluoranthene	-						1 <del>44</del> 1
Benzo(g,h,i)perylene Benzo(j)fluoranthene	191-24-2 205-82-3	-			7.64		7.64
Benzo(k)fluoranthene	207-08-9 218-01-9			-		0.018 0.018	0.018 0.018
Chrysene Dibenzo(a,h)anthracene	53-70-3		1 (444) 1 (444)			0.018	0.018
Fluoranthene Fluorene	206-44-0 86-73-7	1.6 2.5		-17:	1.9 19	140 5300	140 5300
Indeno(1,2,3-c,d)pyrene	193-39-5			**	4.31	0.018	0.018
Naphthalene Phenanthrene	91-20-3 85-01-8	1.4 (a) 1.5	/ <u>122</u> /	<del></del>	13 3.6	1 <del>22</del> 1	13 1.5
Pyrene Total Benzofluoranthenes (b,j,k)	129-00-0	0.24			0.3	4000	4000
Total HPAH	725	122	- (Appli)		100	AMACA.	124
Total LPAH Total PAH	-	-FT				-	\ <del>-</del>
Polychlorinated Biphenyls (PCBs) (ug/L)	10071440		T and	272	(Mari	1221	
Aroclor 1221	12674-11-2 11104-28-2		(20) 				-
Aroclor 1232 Aroclor 1242	11141-16-5 53469-21-9						-
Aroclor 1242 Aroclor 1248	12672-29-6	7.75					
Aroclor 1254 Aroclor 1260	11097-69-1 11096-82-5				-		
Aroclor 1262	37324-23-5	1999		: (==)			(200)
Aroclor 1268 Semivolatile Organic Carbons (SVOCs) (ug/L)	11100-14-4	155		1550		<b>m</b> s	20
1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene	95-94-3 120-82-1	129 5.4 (a)	(840)	***	3 30	1.1 70	1.1 70
1,2-Dichlorobenzene	95-50-1	42 (a)	1970)		14	1300	1300
1,3-Dichlorobenzene 1,4-Dichlorobenzene	541-73-1 106-46-7	28.5 19.9	(2000) (2000)		38 9.4	960 190	960 190
2,2'-Oxybis (1-chloropropane)	108-60-1	- 12	1607	-		65000	65000 1.2
2,3,4,6-Tetrachlorophenol 2,4,5-Trichlorophenol	58-90-2 95-95-4	1.2 12	1771 1784		1.2	1 <del></del> 1	12
2,4,6-Trichlorophenol 2,4-Dichlorophenol	88-06-2 120-83-2	61 11	(Aug.		4.9 11	2.4 290	2.4 290
2,4-Dimethylphenol	105-67-9		0750	<del></del> -	100	850	850
2,4-Dinitrophenol 2,4-Dinitrotoluene	51-28-5 121-14-2	48.5 44	(See		19 44	5300 3.4	5300 3.4
2,6-Dinitrotoluene 2-Chloronaphthalene	606-20-2 91-58-7	81	(BS)		81 0.396	1600	81 1600
2-Chlorophenol	95-57-8	265	·		24	150	150
2-Methylphenol (o-Cresol) 2-Nitroaniline	95-48-7 88-74-4	1020	120		67	1221	1020
2-Nitrophenol 3,3'-Dichlorobenzidine	88-75-5 91-94-1	2940 73		_	 4.5	0.028	2940 0.028
3-Methylphenol & 4-Methylphenol (m&p-Cresol)	1319-77-3	- Jan		-			1 444
3-Methylphenol (m-Cresol) 3-Nitroaniline	108-39-4 99-09-2		- 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1	<u></u>	62	1200	62 
4-Bromophenyl-phenyl ether	101-55-3	1.5			1.5		1.5 34.8
4-Chloro-3-methylphenol 4-Chloroaniline	59-50-7 106-47-8	232	(25)		34.8 232	(144)	232
4-Methylphenol (p-Cresol) 4-Nitroaniline	106-44-5 100-01-6	543	,		25		543 
4-Nitrophenol	100-02-7	71.7	(144)		60	CAMP	71.7
Acetophenone Aniline	98-86-2 62-53-3	2.2	(455) (55)		4.1	(122)	2.2
Atrazine Benzaldehyde	1912-24-9 100-52-7	1.8	( <del></del> )			100	1.8
Benzidine	92-87-5	3.9	100			(22)	3.9
Benzoic acid Benzyl alcohol	65-85-0 100-51-6	42 8.6		-	8.6	1550	42 8.6
Biphenyl (1,1'-Biphenyl)	92-52-4	14	(244) (447)			1221	14
bis(2-Chloroethoxy)methane bis(2-Chloroethyl)ether	111-91-1 111-44-4		100	-	19000	0.53	0.53
bis(2-Ethylhexyl)phthalate Butylbenzyl phthalate	117-81-7 85-68-7	16 29.4			0.3	2.2 1900	2.2 1900
Caprolactam	105-60-2	- M	(ES)		22	1220	
Dibenzofuran Diethyl phthalate	132-64-9 84-66-2	65 75.9			4 110	44000	65 44000
Dimethyl phthalate	131-11-3	580	(200)			1100000	1100000 4500
Di-n-butyl phthalate Dinitro-o-cresol (4,6-Dinitro-2-methylphenol)	84-74-2 534-52-1	3.4	(20) (20)	-	9.7 23	4500 280	280
Di-n-octyl phthalate Hexachlorobenzene	117-84-0 118-74-1	22 0.0003			30 0.0003	0.00029	22 0.00029
Hexachlorocyclopentadiene	77-47-4	0.07	1887	122	77	1100	1100
Hexachloroethane Isophorone	67-72-1 78-59-1	9.4 129			8 920	3.3 960	3.3 960
Nitrobenzene	98-95-3 62-75-9	66.8 330000	(ALC)		220	690	690 330000
n-Nitrosodimethylamine		330000 120		-		0.51	0.51
n-Nitrosodi-n-propylamine	621-64-7						
THE RESERVE THE PARTY OF THE PA	86-30-6 87-86-5	33000 7.9	7.9	 13	4	6 3	6 3

Analyte	CAS Number	EPA Region 3 BTAG Marine Water Screening Benchmarks EPA. 2006	National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CCC (chronic) <sup>1</sup> EPA, 2013	National Recommended Water Quality Criteria - Aquatic Life Criteria - Saltwater CMC (acute) <sup>1</sup> EPA. 2013	EPA Region 5 RCRA - Ecological Screening Levels - Water EPA. 2003	National Recommended Water Quality Criteria - Human Health for the Consumption of Organisms EPA, 2013	Initial PRGs Used for Data Screening
Volatile Organic Carbons (VOCs) (ug/L)	CAS (Vallibe)	2171, 2000	2111, 2010	1111/2020	2171,2505	2719 2020	TOI Data Screening
1,1,1-Trichloroethane	71-55-6	312		**	76	_	312
1,1,1,2-Tetrachloroethane	630-20-6					-	
1,1,2,2-Tetrachloroethane	79-34-5	90.2	0.000		380	4	4
1,1,2-Trichloroethane	79-00-5	550	124	N <u>24</u> 6	500	16	16
1,1,2-Trichlorotrifluoroethane (Freon 113)	76-13-1		-				
1,1-Dichloroethane	75-34-3	47	SI <del>NE</del> S		47	1==1	47
1,1-Dichloroethene	75-35-4	2240	0440		65	7100	7100
1,2,3-Trichlorobenzene	87-61-6	8	158165		122	15050	8
1,2,3-Trichloropropane	96-18-4		(58)			1550	550
1,2,4-Trimethylbenzene	95-63-6	19		**			19
1,2-Dibromo-3-chloropropane	96-12-8	-	0==0	-			1441
1,2-Dichloroethane	107-06-2	1130	1240	_	910	37	37
1,2-Dichloroethene, cis-	156-59-2	-	(68)	-			<del></del>
1,2-Dichloroethene, trans-	156-60-5	970	10 <b>44</b> 1		970	10000	10000
1,2-Dichloropropane	78-87-5	2400	(1940)		360	15	15
1,3,5-Trimethylbenzene (Mesitylene)	108-67-8	71	/100		-22		71
1,3-Dichloropropane	142-28-9	=	(55)	***	No.	(50)	ms
1,3-Dichloropropene, cis-	10061-01-5	-	OME	**	366	21	21
1,3-Dichloropropene, trans-	10061-02-6	-	3 <b>44</b> 2	-		21	21
1,4-Dichloro-2-butene, trans-	110-57-6	622	1967	122	192	(1220)	1997
1,4-Dioxane	123-91-1	-	(57)	-	22000		22000
2-Butanone (MEK)	78-93-3	14000	0##0		2200	1	14000
2-Chlorotoluene	95-49-8	-	5 <b>44</b> 0		CHH.	(m)	1441
2-Hexanone (Methyl butyl ketone)	591-78-6	99	1999	122	99	12290	99
4-Chlorotoluene	106-43-4	-	(572)	-	157		
4-Isopropyltoluene (4-Cymene)	99-87-6	85	0.00				85
Acetone	67-64-1	564000	(1 <b>44</b> )		1700		564000
Acrolein	107-02-8	0.55	(86)	123	0.19	1221	0.55
Acrylonitrile	107-13-1	581	(55)		66	11500	581
Benzene	71-43-2	110 (a)		***	114	51	51
Bromobenzene	108-86-1		(144)	***		(44)	1641
Bromochloromethane	74-97-5		(56)	123	122	1221	U-
Bromodichloromethane	75-27-4	-	- 550	-		17	17
Bromoform (Tribromomethane)	75-25-2	640	G##5		230	140	140
Bromomethane (Methyl bromide)	74-83-9	120	(See		16	1500	1500
Carbon disulfide	75-15-0	0.92	1960		15	16	0.92
Carbon tetrachloride (Tetrachloromethane)	56-23-5	1500	1576)	-	240	1.6	1.6
Chlorobenzene	108-90-7	25 (a)	COMMO		47	1600	1600
Chloroethane Chloroform	75-00-3 67-66-3	 815	1257		140	470	470
The state of the s	74-87-3	2700				4/0	2700
Chloromethane Cyclohexane	110-82-7						
Dibromochloromethane	124-48-1	-				13	13
Dibromomethane	74-95-3		12.2		122	15	- 15
Dichlorodifluoromethane	75-71-8	-					-
Dichloromethane (Methylene chloride)	75-09-2	2560	(and)		940	590	590
Ethylbenzene	100-41-4	25 (a)			14	2100	2100
Ethylene dibromide (1,2-Dibromoethane)	106-93-4		- 122		222		_
Hexachlorobutadiene (Hexachloro-1,3-butadiene)	87-68-3	0.3	( <u>9</u> 78)		0.053	18	18
Isopropylbenzene (Cumene)	98-82-8	2.6	1000	_		-	2.6
Methyl acetate	79-20-9	=-	28 <b>44</b> 0		544	7440	-
Methyl iodide (lodomethane)	74-88-4		160		100	1221	
Methyl isobutyl ketone (4-Methyl-2-pentanone or (MIBK))	108-10-1	123000	( <del>110</del> )	_	170	1500	123000
Methyl tert-butyl ether (MTBE)	1634-04-4	11070				-	11070
n-Butylbenzene	104-51-8	-	S1846		SHA		<b>1</b> =1
n-Propylbenzene	103-65-1	128	(Section)	www.	122	1220	128
o-Xylene	95-47-6		(55)	I MORE	- 5.5	1980	ms.
sec-Butylbenzene	135-98-8				**	-	
Styrene	100-42-5	910	S1846		32	THE STATE OF THE S	910
tert-Butylbenzene	98-06-6	E	(202)			1920	_
Tetrachloroethene (PCE)	127-18-4	45	(55)		45	3.3	3.3
Toluene	108-88-3	215 (a)			253	15000	15000
Total xylene (reported, not calculated)	1330-20-7						1440
Total Xylene	-	19	/A24	and the second	27	1 <u>20</u> 0	19
Trichloroethene (TCE)	79-01-6	21	0570	-	47	30	30
Trichlorofluoromethane (Fluorotrichloromethane)	75-69-4	-	S <del>100</del> 0			1000	н
			r				

Vinyl acetate Vinyl chloride

Compounds frequently associated with MGP-operations

1 = Criteria for metals and methyl mercury are expressed in terms of the dissolved metal in the water column.

(a) = This is a Canadian Water Quality Guideline value and refers to the total concentration in an unfiltered sample.

BTAG = Biological Technical Assistance Group

CAS = Chemical Abstract Services

CCC = Criterion Continuous Concentration

CMC = Criterion Maximum Concentration

EPA = U.S. Environmental Protection Agency

HPAH = high molecular weight PAH

108-05-4

EPA = U.S. Environmental Protection Agency
HPAH = high molecular weight PAH
LPAH = low molecular weight PAH
L = liter
mg = miligram
MGP = manufactured gas plant
ng = nanogram
PRG = Preliminary Remediation Goal
RCRA = Rsource Conservation and Recovery Act
RSL = rezional screening level

RSL = regional screening level ug = microgram

EPA, 2003. EPA Region 5 Resource Conservation Recovery Act (RCRA) Ecological Screening Levels. August 22, 2003.

EPA, 2006. EPA Region 3 Biological Technical Assistance Group (BTAG) Screening Benchmarks. Marine Sediment Benchmarks. July 2006.

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EPA, 2013a. National Recommended Water Quality Criteria. Updated August 22, 2013. Available from: http://water.epa.gov/scitech/swguidance/standards/criteria/current/index.cfm#altable.

### Table 7-1 - Summary of Data Quality Review for Existing Site Data Bremerton Gas Works Site Bremerton, Waahington

				Study/Media			
	2008 E&E Targeted Brownfields (E&E 2008)	2010 E&E Removal Action (AnchorQEA 2011)	1995 Ecology (Ecology 1995)	2007 Geoengineers (Geoengineers 2007a)	2008 E&E Targeted Brownfields (E&E 2008)	2007 Geoengineers (Geoengineers 2007a)	2008 E&E Targeted Brownfields (E&E 2008)
	Sediment	Sediment	Soil	Soil	Soil	Groundwater	Groundwater
Work Plan Documentation							
Work Plan (SAP/QAPP)	Detailed QAPP covering multiple pieces of sampling program (soil, groundwater and sediment). Also includes general sediment sampling SOP and data report.	Site-Specific Sampling Plan (SSSP; not reviewed) approved by EPA, finalized after sampling conducted but in field deviations approved by EPA.	, None	Work Plan, including site-specific SAP and QAPP, dated June 1, 2007	SQAPP dated March 5, 2008	Work Plan, including site-specific SAP and QAPP, dated June 1, 2007	SQAPP dated March 5, 2008
Collection methods and purpose	Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures. Limited for sediment; to determine if GW migration from upland sources is occurring into the Narrows.	Developed under EPA Superfund Tecncial Assessment Respoonse Tear (START). Determining origin of contamination from 12" exposed drain pipe on Sesko property beach.	Surface soil/sediment samples of suspected contamination	Purpose to assess soil quality in potential contaminant source areas. Table of rationale for specific boring/sample locations referenced but not included in final work plan.	Judgmental sampling design to determine presence of contamination in areas of concern. Detailed rationale provided in SQAPP.	Purpose to assess groundwater quality in and downgradient of potential contaminant source areas. Table of rationale for specific boring/sample locations referenced but not included in final work plan.	Judgmental sampling design to determine presence of contamination in areas of concern. Detailed rationale provided in SQAPP.
Sample Location and Collection Methods							
Location method, accuracy, and datum.	Location established with GPS coordinates; accuracy not specified. Actual sampling appear to be close/at QAPP locations. Datum not specified.	Location established with GPS coordinates; accuracy not specified.  Datum not specified.	Sample locations recorded on rough site sketch. No survey information provided.	Locations provided on scaled site map. Location method unknown. No survey information provided.	Locations provided on scaled site map. Location method unknown. No survey information provided. Note: locations of borings SP01 and SP03 apparently switched on site map, based on boring log information and correlation of chemical data with boring log observations.	Locations provided on scaled site map. Location method unknown. No survey information provided.	Locations provided on scaled site map. Location method unknown. No survey information provided. Note: locations of borings SP01 and SP03 apparently switched on site map, based on boring log information and correlation of chemical data with boring log observations.
Sample depths	0-30cm	0-6 inches	Less than 10 inches	up to 45 feet deep	up to 40 feet deep	15-foot long well screens up to 45 feet deep	Monitoring Wells: 10-foot long well screens up to 45 feet deep. Temporary borings: depth not provided.
Collection method and matrix	Surface sediment. Dedicated stainless steel spoon. Collected at low tide from 5 biased locations targeted to evaluate potential for GW migration based on previous analytical and "on-site observations".	Surface sediment. Dedicated stainless steel spoon. Known areas of sediment deposition within the direct vicinity of the 12: drainpipe, collected below average high tide line.	Hand collection of surface soil/sediment samples	Hollow-stem auger drilling with split-spoon sampling.	Hollow-stem auger drilling with split-spoon sampling.	Report states low-flow sampling with peristaltic pump. Questionable for 30-ft deep groundwater samples.	Monitoring wells sampled using low-flow sampling using electric submersible pump. Methods for sampling temporary boreholes not provided.
Sample collection, processing and handling	Homogenized in dedicated stainless steel bowls (VOC cores taken from sampling locations prior to other sediment collection). Data report includes photographs at each sediment station.	Homogenized in dedicated stainless steel bowls (VOC cores taken from sampling locations prior to other sediment collection). Data report includes photographs at each sediment station.	Collection and handling activities not reported.	Soil samples collected from 8 borings at 5-foot intervals and field screened for contamination. 17 samples collected for sample analysis. VOC samples collected by EPA 5035A. Protocols detailed in SAP.	Soil samples collected from 7 borings at 5-foot intervals and field screened for contamination. 48 samples collected for sample analysis. VOC samples collected by EPA 5035A. Protocols detailed in SAP.	Groundwater samples collected from 8 permanent, developed monitoring wells. Processing and handling protocols detailed in SAP.	Groundwater samples collected from 2 permanent, developed monitoring wells and 4 temporary borings. Processing and handling protocols detailed in SAP.
Holding time, preservation, and chain of custody	Detailed in the QAPP. COCs provided in data report. Holding time and preservation discussed in lab data report.	COCs provided in data report. Holding time and preservation discussed in lab data report.	chain of custody not provided. Laboratory case narrative indicates holding times were within recommended limits.	Requirements detailed in SAP and QAPP. Holding times and preservation were met as documented in data report. Chain of custody provided in data report.	Requirements detailed in SAP and QAPP. Holding times and preservation were met as documented in data report. Chain of custody provided in data report.	Requirements detailed in SAP and QAPP. Holding times and preservation were met as documented in data report. Chain of custody provided in data report.	Requirements detailed in SAP and QAPP. Holding times and preservation were met as documented in data report. Chain of custody provided in data report.
Laboratory Analysis							
Analytical methods are standard or USEPA approved	d EPA and NWTPH methods. TPH-Dx, TPH-Dx, VOC, SVOC, TAL metals.	EPA methods. VOC by 8260, SVOC by 8270, static sheen test.	EPA Methods.  Metals - EPA200.7, EPA270.2, EPA206.2, EPA279.2, EPA245.5  PAHs - Manchester Modification of SW8270	EPA and NWTPH Methods. TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA -8260B SVOCs - EPA 8270 SIM PCBs - EPA 8082 PP metals/chromiumVI - EPA 6000/7000 series TBT - Krone (GC/MS)	TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA 8250B SVOCs - EPA 8270C TAL metals - EPA 6000/7000 series	EPA and NWTPH Methods. TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA -8260B SVOCs - EPA 8270 SIM PCBs - EPA 8082 PP metals/chromiumVI - EPA 6000/7000 series	EPA and NWTPH Methods. TPH - Ecology NWTPH-Gx and NWTPH-Dx VOCs - EPA 8260B SVOCs - EPA 8270C TAL metals - EPA 6000/7000 series
Detection limits and qualifiers determined based on USEPA guidance	Yes. Detailed in the QAPP. Qualifier identified in laboratory data report.	Yes. Qualifier identified in laboratory data rreport.	summarized in QA narrative in laboratory data report.	Yes. Detailed in QAPP. Qualifiers identified in laboratory data report.	Yes. Detailed in QAPP. Qualifiers identified in laboratory data report.	Yes. Detailed in QAPP. Qualifiers identified in laboratory data report.	Yes. Detailed in QAPP. Qualifiers identified in laboratory data report.
Measurement instruments and calibration procedures	Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures.	Some detail provided in data validation memo.	Some detail provided in QA narrative in laboratory data report.	Yes. Detailed in QAPP.	Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures.	Yes. Detailed in QAPP.	Detailed in QAPP. Sampling under EPA Brownfields management, follows EPA procedures.
Quality Control and Data Validation Field/Lab quality control samples (duplicates, blanks)	Field rinsate and trip blanks (no issues in sediment samples) MS/MSD, serial dilution, internal standards.	Field trip blank.	MS/MSD, LCS	Field duplicate; method blanks, calibration blanks, sample blanks, MS/MSD, and LCS.	Laboratory blanks, rinsate blanks, trip blanks, MS/MSD.	Field duplicate, rinseate blank, and trip blanks; method blanks, calibration blanks, sample blanks, MS/MSD, and LCS.	Laboratory blanks, rinsate blanks, trip blanks, MS/MSD.
Analytical chemistry data must have been validated and qualified consistent with EPA functional guidelines	Data validation conducted. Data validation memo included as Appendix to data report. Procedures also detailed in QAPP.	Data validation conducted. Data validation memo included as Appendix to data report.	QA summary by lab. Compounds with low matrix spike recoveries rejected or "J" qualified.	QA summary by lab.	QA/QC review and data validation documented in data report.	QA summary by lab.	QA/QC review and data validation documented in data report.
Laboratory data reports	Level II Data Package Available.	Level II Data Package Available.	Partial Level II Data Package Available.	Level II Data Package Available.	Level II Data Package Available.	Level II Data Package Available.	Level II Data Package Available.

COC = chemical of concern
EPA = U.S. Environmental Protection Agency
GC/MS = gas chromatography-mass spectrometry

LCS = laboratory control sample MS/MSD = matrix spike/matrix spike duplicate NWTPH = Northwest total petroleum hydrocarbon

PAH = polycyclic aromatic hydrocarbon PCB = polychlorinated biphenyl PP = priority pollutant QA = quality assurance QAP = quality assurance
QAPP = Quality Assurance Project Plan
QC = quality control
SAP = Sampling and Analysis Plan
SOP = standard operating procedure
SQAPP = SAP/QAPP SVOC = semivolatile organic compound
TAL = target analyte list
TBT = tributyltin
TPH = total petroleum hydrocarbons VOC = volatile organic compound

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Ecology & Environment, Inc. (E&E), 2008, Final Bremerton Gasworks Targeted Brownfields Assessment Sampling and Quality Assurance Project Plan, Prepared by E&E for EPA, March 5, 2008.

Anchor QEA, 2011, Final Completion Report: Former Bremerton MGP Site, Incident Action and Time Critical Removal Action, Prepared for U.S. Coast Guard Sector Puget Sound Incident Management Division on behalf of Cascade Natural Gas Corporation, January 2011.

Bremerton Gas Works Site Bremerton, WA

			Study/Media			_
	2010 and 2012 ENVVEST	2005, 2007 NOAA Mussel Watch @ station SIWF	2001 303d Ecology Clam Crab	2008-2009 PSAMP - Spatial/Temporal - Central Sound	1989-2013 PSAMP Long term/ temporal	2009 - PSAMP Urban Waters Initiative - Bainbridge Basin
	Mussel tissue. Data from 11 locations in Dyes Inlet and Sinclair Inlet considered for regional information.	Mussel Tissue. Data from 1 location in Sinclair Inlet considered for regional information.	Clam and crab tissue. Data from 3 locations in Dyes Inlet considered for regional information.	Sediment. Data from 11 locations in Dyes Inlet and Sinclair Inlet considered for regional information.	Sediment. Data from 1 location in Sinclair Inlet considered for regional information.	Sediment. Data from 18 locations in Dyes Inlet and Sinclair Inlet considered for regional information.
Work Plan Documentation			_	_		
Work Plan (SAP/QAPP)	Detailed SAP/QAPP developed with EPA and Ecology under the coorperative Environmental Investment (ENVVEST) program (Johnston et al. 2009; 2010).	Detailed SAP/QAPP developed under NOAA National Status and Trends Program (NOAA 1993 and 2006).	Ecology (2001) QAPP. Results summarized in the 2002 data report and queried from EIM.	Detailed programattic QAPP (2009) developed cooperatively with State and Federal agencies. Event specific addenda (2010, 2011, 2012).	Detailed programattic QAPP (2009) developed t-cooperatively with State and Federal agencies. Event specific addenda (2010, 2011, 2012).	Detailed programattic QAPP (2009) developed cooperatively with State and Federal agencies. Event specific addenda (2010, 2011, 2012).
Collection methods, purpose and representativeness	Hand collection of blue mussels (Mytilus spp.) via boat or from shore. Shucked, s whole organism. Methods follow NOAA protocol. Location control details provided.	Hand collection of blue mussels (Myzilus spp.) via boat or from shore. Shucked, whole organism. Methods follow NOAA protocol.	Hand collection of male cancer crab tissue (Cancer gracilis) via crab pots (though Dungeness and Blue crabs targeted but none found); native and Japanese little neck clam tissue via hand digging (Protothaca staminea and Tapes japonica).	0.1 m2 modified stainless steel van Veen, lowered via cable to open upon sediment contact. Targeted fine grained sediment, sample rejected in field if not fine-grained dominant during in-field visual inspection.	0.1 m2 modified stainless steel van Veen, lowered via cable to open upon sediment contact. Targeted fine grained sediment, sample rejected in field if not fine-grained dominant during in-field visual inspection.	0.1 m2 modified stainless steel van Veen, lowered via cable to open upon sediment contact. Targeted fine grained sediment, sample rejected in field if not fine-grained dominant during in-field visual inspection.
Sample Location and Collection Methods						
Location method, accuracy and datum	Location established with GPS; accuracy not specified. Table provided with coordinates. Datum not specified.	Location established with GPS. Accuracy and datum not specified.	Location established with GPS, accuracy not specified. Table provided with coordinates. Datum is NAD 83.	Location established with differential GPS. with expected accuracy of better than 3 meters. Table provided with coordinates. Datum is NAD 83.	Location established with differential GPS. with expected accuracy of better than 3 meters. Table provided with coordinates. Datum is NAD 83.	Location established with differential GPS. with expected accuracy of better than 3 meters. Table provided with coordinates. Datum is NAD 83.
Sample depths	Above MLLW - on rocks, piling, cabling, piers.	Detailed in NOAA (1993) SAP. Depends on station, some shoreline, some underwater.	Crabs: via pots on surface Clams: via hand digging within 100 sq ft of beach.	Top 2-3cm.	Top 2-3cm.	Top 2-3cm.
Sample collection, processing and handling	Field - Hand harvest, cut byssus threads with knife; hand brush off debris; 1-3 replicates per stations (reps within 150' radius of station loc; 30-50 mussels per replicate. Hand delivery to lab.  Lab - kept at -20C until measured and shucked with ceramic knife; rinsed with DI, composite by replicate then by station using Ti blender.	Field - Detailed in NOAA (1993) SAP. In general, some stations hand collection or with rake, some with bivalve dredge.  Lab - shell size and volume determined; shucked; homogenized using stainless steel blender with titanium blades. Chemically dried using hydromatrix.	Detailed in SAP.Crabs: Muscle tissue (no organs or shell). Clams: Non depurated. Both crabs and clams samples homogenized in stainless steel blender.	Field - stainless steel spoon from each grab; grabs composited into stainless steel bucket; salinity and sediment temp measured.	Field - stainless steel spoon from each grab; grabs composited into stainless steel bucket; salinity and sediment temp measured.	Field - stainless steel spoon from each grab; grabs composited into stainless steel bucket; salinity and sediment temp measured.
Holding time, preservation, and chain of custody	Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report.	Procedures detailed in NOAA (1993) SAP. Actual COCs not available.	Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report.	Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report.	Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report.	Requirements detailed in the QAPP. Holding times and preservation were met as documented in the data report. Chain of custody provided in the data report.
Laboratory Analysis						·
Analytical methods are standard or EPA approved	Total Hg - EPA 7473m (EPA 1631 rev E in QAPP). Battelle SOPS for other metals and PCB congeners, PAHs - GC/MS Battelle SOP -015. Standard analytical methods. Lipids, moisture, C and N isotopes, trace metals, Hg, isotopes, 20 NS&T PCB congeners, parent and alkylated PAH.	Lipids, moisture, C and N isotopes, trace metals, Hg. isotopes, 20 NS&T PCB congeners, parent and alkylated PAH. Detailed in specific analytical methods reports. Standard anlytical methods.	Lipid, and imony, SVOCs, PAHs. USEPA and PSEP standard anlytical methods.	Grain size, TOC, metals, pesticides, chlorobenzenes, PAHs, phenolics, phthalates, PCBs, PBDEs, bPA, triclosan, and other misc. including HCBD, dibenzofuran, carbazole and tin. EPA and PSEP standard anlytical methods.	USEPA and PSEP standard anlytical methods.	USEPA and PSEP standard anlytical methods.
Detection limits and qualifiers determined based on EPA guidance	Yes. Detailed in QAPP and summarized in QA/QC narrative in data report.	Yes. Detailed in QAPP and summarized in QA/QC narrative in data report.	Yes. Detailed in QAPP and summarized in QA/QC narrative in data report.	Yes. Detailed in QAPP and summarized in QA/QC narrative in data report.	Yes. Detailed in QAPP and summarized in QA/QC narrative in data report.	Yes. Detailed in QAPP and summarized in QA/QC narrative in data report.
Measurement instruments and calibration procedures	Detailed in QAPP.	Detailed in QAPP.	Detailed in QAPP.	Detailed in QAPP.	Detailed in QAPP.	Detailed in QAPP.
Quality Control and Data Validation			1			T
Field/Lab quality control samples (duplicates, blanks	) B, BS, MS/MSD, LD, reference material.	B, BS, MS/MSD, LD, reference material.	Blank, MS/MSD.	Blind field split replicates, field blanks; lab replicates, MS/MSD, lab control, MB, reference material.	, Blind field split replicates, field blanks; lab replicates, MS/MSD, lab control, MB, reference material.	Blind field split replicates, field blanks; lab replicates, MS/MSD, lab control, MB, reference material.
Analytical chemistry data must have been validated and qualified consistent with EPA functional guidelines	Data validation conducted. Details in case narratives.	Not available online.	Data validation conducted. Details in case narratives.	Data validation conducted. Details in case narratives.	Data validation conducted. Details in case narratives.	Data validation conducted. Details in case narratives.
Laboratory data reports	Level II Data Package Available.	Not available online.	Case narrative text only.	Level II Data Package Available.	Only case narratives available through 2000. Online archives incomplete.	Level II Data Package Available.

Notes: References: B = Blank 1989-2013 PSAMP bPA = Bisphenol A 2008-2009 PSAMP BS = Blank spike 2009 PSAMP COCs = chemical of concerns EPA = U.S. Environmental Protection Agency HCBD = Hexachlorobutadiene 2010 and 2012 ENVVEST (Johnston 2010 and Brandenberger 2012) GPS = global positioning system Johnston et al. 2009: 2010 LCS = Laboratory control sample MB = Method blank MS/MSD = Matrix spike/matrix spike duplicate MLLW = Mean lower-low water 2005, 2007 NOAA Mussel Watch NOAA = National Oceanic and Atmospheric

2001 303d Ecology Clam Crab

Striplin, P.L., 1988. Puget Sound Ambient Monitoring Program: Marine Sediment Quality Implementation Plan. Washington State Department of Ecology, Olympia, Washington. 57 pp. www.ecy.wa.gov/biblio/88e37.html. Also see QAPP addendum PSAMP (2009, 2010, 2011, and 2012). PSAMP. 2009. Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program: Sediment Monitoring Component. August 2009. Publication No. 09-03-121

PSAMP. 2010 Addendum to Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program: Sediment Monitoring Component. August 2010. Publication No. 09-03-121-Addendum1

PSAMP. 2011 Addendum to Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program: Sediment Monitoring Component. August 2010. Publication No. 09-03-121-Addendum2
PSAMP. 2012 Addendum to Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program/Urban Waters Initiative: Sediment Monitoring in the San Juan Islands and Port Gardner/ Everett Harbor. December 2011. Publication No. 09-03-121-Addendum3

PSAMP. 2012 Addendum to Quality Assurance Project Plan. The Puget Sound Assessment and Monitoring Program/Urban Waters Initiative: Sediment Monitoring in the San Juan Islands and Port Gardnery Everett Harbor. December 2011. Publication No. 09-03-121-Addendums
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Administration

PAHs = polycyclic aromatic hydrocarbons

PBDE = Polybrominated diphenyl ether PCBs = polychlorinated biphenyls PSBP = Puget Sound Estuary Program QAPP = Quality Assurance Project Plan QA/QC = quality assurance/quality control SAP = Sampling and Analysis Plan SVOC = semivolatile organic compound TAL = Target analyte list TOC = Total organic carbon TPH = total petroleum hydrocarbons VOC = volatile organic compound

### **Table 7-3 - Statistical Summary of Soil Data**Bremerton Gas Works Site

Chemical Group	Chemical Constituent	Number of Locations	Number of Samples	Number of Detections	Maximum Detected Concentration (mg/kg)	Minimum Detected Concentration (mg/kg)	Soil Initial PRG (mg/kg)	Detected Concentrations Exceeding the Initial PRG	Results with Reporting Limit Concentrations that Exceed the Initial Soil PRG	Puget Sound Background Metals Concentration (mg/kg) <sup>1</sup>	Number of Detected Concentrations Exceeding Puget Sound Background	Number of Non-Detect Results with Reporting Limit Concentrations that Exceed Puget Sound Background
	Gasoline Range Hydrocarbons	15	59	11	645	5						
TPH	Diesel Range Hydrocarbons	15	58	10	36000	17.1						
	Oil Range Hydrocarbons	15	58	11	29000	18						
	Aluminum	7	42	42	24100	5780	77000			32600		
	Antimony	13	31	2	1.2	0.8	0.27	2	29	5		12
	Arsenic	15	59	59	48.4	0.5	0.61	55		7	2	
	Barium	7	42	42	120	23.9	330			255		
	Beryllium	15	59	42	0.5	0.1	21			0.6		7
	Cadmium	15	59	34	1.6	0.2	0.36	21	25	1	4	
	Calcium	7	42	42	21300	1620						
	Chromium (Total)	15	59	59	60.8	14.6	26	32		48	11	
	Chromium (VI)	8	17	0	NA	NA	0.29		17			
	Cobalt	7	42	42	19	3.3	13	13		11	15	
	Copper	15	59	59	79.1	8	28	18		36	17	
Metals	Iron	7	42	42	47800	9570	55000			36100	3	
ivictals	Lead	15	59	57	246	0.6	11	7		24	6	
	Magnesium	7	42	42	14900	1380						
	Manganese	7	42	42	824	170	220	26		1200		
	Mercury	15	59	14	1.62	0.1	10			0.07	14	45
	Nickel	15	59	59	66.3	21.2	38	27		48	17	
	Potassium	7	42	42	2000	233						
	Selenium	15	59	0	NA	NA	0.52		57	0.78		42
	Silver	15	59	0	NA	NA	4.2			0.61		46
	Sodium	7	42	42	565	120						
	Thallium	15	59	34	5.7	1.1	0.78	34	8			
	Vanadium	7	42	42	86	20.7	7.8	42		45	17	
	Zinc	15	59	59	376	18.9	46	23		85	5	
	Acenaphthene	18	60	19	31.2	0.0012	3400					
	Acenaphthylene	23	61	23	460	0.00091	682					
	Anthracene	20	61	24	274	0.0012	17000					
	Benzo(g,h,i)perylene	19	61	46	79	0.00071	119					
	Dibenzofuran	15	59	4	0.37	0.017	78		2			
PAHs	Fluoranthene	22	61	32	572	0.00068	2300					
	Fluorene	20	61	25	404	0.0007	2300					
	Phenanthrene	24	61	39	1490	0.00061	45.7	6				
	Pyrene	21	61	38	913	0.0006	1700					
	1-Methylnaphthalene	12	17	10	615	0.0144	16	5				
	2-Methylnaphthalene	13	17	10	978	0.0158	230	2				
	Naphthalene	10	12	11	953	0.00047	3.6	4				
T	Benz(a)anthracene	18	61	29	113	0.0011	0.15	15	2			
L	Benzo(a) pyrene	17	61	40	116	0.00053	0.015	21				
	Benzo(b)fluoranthene	17	61	29	57.4	0.00085	0.15	16	1			
	Benzo(k)fluoranthene	17	61	36	60.6	0.00056	1.5	10				
cPAHs	Chrysene	17	61	35	146	0.00067	15	6				
	Dibenzo(a,h)anthracene	17	61	36	22.8	0.0008	0.015	16	3			
	Indeno(1,2,3-cd)pyrene	17	61	44	58.5	0.00066	0.15	15	1			
	Total cPAHs TEQ (ND = 0)	17	61	50	149	0.000066	0.015	21				
	Total cPAHs TEQ (ND = 1/2 RDL)	17	61	50	149	0.000842	0.015	22				

### **Table 7-3 - Statistical Summary of Soil Data**Bremerton Gas Works Site

					Maximum	Minimum		Detected	Results with Reporting	Puget Sound	Number of Detected	Number of Non-Detect Results
					Detected	Detected		Concentrations	Limit Concentrations	Background Metals	Concentrations	with Reporting Limit
Chemical		Number of	Number of	Number of	Concentration	Concentration	Soil Initial PRG	Exceeding the	that Exceed the Initial	Concentration	Exceeding Puget	Concentrations that Exceed
Group	Chemical Constituent	Locations	Samples	Detections	(mg/kg)	(mg/kg)	(mg/kg)	Initial PRG	Soil PRG	(mg/kg) <sup>1</sup>	Sound Background	Puget Sound Background
	1,1'-Biphenyl	7	42	5	0.98	0.014	51					
	1,2,4,5-Tetrachlorobenzene	7	42	0	NA	NA	18					
	1,2,4-Trichlorobenzene	15	59	2	0.00023	0.00014	22					
	1,2-Dichlorobenzene	1	1	0	NA	NA	1900					
	1,3-Dichlorobenzene	7	40	0	NA	NA	37.7					
	1,4-Dichlorobenzene	1	1	0	NA	NA	2.4					
	1,4-Dioxane	7	42	0	NA	NA	4.6		2			
	2,3,4,6-Tetrachlorophenol	7	42	0	NA	NA	1800					
	2,4,5-Trichlorophenol	15	59	0	NA	NA	6100					
	2,4,6-Trichlorophenol	15	59	0	NA	NA	44		2			
	2,4-Dichlorophenol	15	59	0	NA	NA	180					
	2,4-Dimethylphenol	15	59	1	0.031	0.031	1200					
	2,4-Dinitrophenol	15	59	0	NA	NA	120		2			
	2-Chloronaphthalene	15	59	0	NA	NA	6300					
	2-Chlorophenol	15	59	0	NA	NA	390					
	2-Methylphenol	8	17	0	NA	NA	3100					
	2-Nitroaniline	15	59	0	NA	NA	610					
	2-Nitrophenol	15	59	0	NA	NA	1.6		8			
	3 & 4 Methylphenol	8	17	0	NA	NA						
	3,3'-Dichlorobenzidine	15	59	0	NA	NA	1.1		17			
	3-Nitroaniline	15	59	0	NA	NA	3.16		7			
1 1	4,6-Dinitro-2-methylphenol	15	59	0	NA	NA	4.9		7			
1 1	4-Bromophenyl phenyl ether	15	59	0	NA	NA						
1 1	4-Chloro-3-methylphenol	15	59	0	NA	NA	6100					
Other	4-Chloroaniline	15	59	0	NA	NA	2.4		8			
SVOCs	4-Chlorophenyl phenyl ether	15	59	0	NA	NA						
	4-Methylphenol	7	42	0	NA	NA	6100					
1	4-Nitroaniline	15	59	0	NA	NA	24		2			
1	4-Nitrophenol	15	59	0	NA	NA	5.12		7			
1	Acetophenone	7	42	2	1.5	0.03	7800					
	Aniline	8	17	0	NA	NA	85		2			
İ	Atrazine	7	42	0	NA	NA	2.1					
İ	Benzaldehyde	7	42	0	NA	NA	7800					
İ	Benzidine	7	42	0	NA	NA	0.0005		42			
İ	Benzoic acid	8	17	0	NA	NA	240000					
İ	Benzyl alcohol	8	17	0	NA	NA	6100					
İ	Benzyl butyl phthalate	15	59	5	0.029	0.015	260					
İ	Bis(2-chloro-1-methylethyl) ether	15	59	0	NA	NA	4.6		4			
	Bis(2-chloroethoxy)methane	15	59	0	NA	NA	180					
	Bis(2-chloroethyl) ether	15	59	0	NA	NA	0.21		17			
	Bis(2-ethylhexyl) phthalate	15	59	39	0.29	0.069	35		2			
	Caprolactam	7	42	1	0.015	0.015	30000					
	Carbazole	15	59	5	0.49	0.019						
	Dibenzofuran	15	59	4	0.37	0.017	78		2			
	Diethyl phthalate	15	59	0	NA	NA	49000		_			
	Dimethyl phthalate	15	59	0	NA NA	NA NA	734					
	Di-n-butyl phthalate	15	59	3	0.016	0.013	6100					
	Di-n-octyl phthalate	15	59	0	NA	NA NA	610					
	Hexachlorobenzene	15	59	0	NA NA	NA NA	0.3		17			
	Hexachlorobutadiene	1	1	0	NA NA	NA NA	6.2					
	Treademorphatamente		<u> </u>		14/3	14/3	0.2					

### **Table 7-3 - Statistical Summary of Soil Data**Bremerton Gas Works Site

Chemical Group	Chemical Constituent	Number of	Number of Samples	Number of Detections	Maximum Detected Concentration (mg/kg)	Minimum Detected Concentration (mg/kg)	Soil Initial PRG (mg/kg)	Detected Concentrations Exceeding the Initial PRG	Results with Reporting Limit Concentrations that Exceed the Initial Soil PRG	Puget Sound Background Metals Concentration (mg/kg) <sup>1</sup>	Number of Detected Concentrations Exceeding Puget Sound Background	Number of Non-Detect Results with Reporting Limit Concentrations that Exceed Puget Sound Background
Стоир		_	-	l l				IIIIIIIII FRG	John Rd	(IIIg/Ng)	J Sound Background	ruget Souria Backgrouna
-	Hexachlorocyclopentadiene	15	59	0	NA NA	NA NA	370					
	Hexachloroethane	15	56	0	NA 6.2	NA 6.2	12		2			
	Isophorone	15	59	1	6.3	6.3	510					
Other	Nitrobenzene	8	17	0	NA NA	NA NA	4.8		4			
L	N-Nitrosodimethylamine	7	42	0	NA NA	NA NA	0.0023		42			
SVOCs	N-Nitroso-di-n-propylamine	15	59	0	NA NA	NA NA	0.069		17			
(continued)	N-Nitrosodiphenylamine	15	59	0	NA 0.0026	NA 0.00004	99		1			
	Pentachlorophenol	15	59	3	0.0036	0.00081	0.89		10			
	Phenol	15	59	6	0.1	0.023	18000					
	2,4-Dinitrotoluene	8	17	0	NA NA	NA	1.6		8			
	2,6-Dinitrotoluene	8	17	0	NA NA	NA NA	0.0328		17			
-	1,1,1,2-Tetrachloroethane	15	59	0	NA	NA	1.9		2			
-	1,1,1-Trichloroethane	15	59	0	NA NA	NA NA	8700					
-	1,1,2 - Trichlorotrifluoroethane	7	42	0	NA	NA	43000		_			
-	1,1,2,2-Tetrachloroethane	15	59	0	NA	NA	0.56		3			
	1,1,2-Trichloroethane	15	59	0	NA	NA	1.1		2			
	1,1-Dichloroethane	15	59	0	NA	NA	3.3		2			
	1,1-Dichloroethene	15	57	0	NA	NA	240					
	1,1-Dichloropropene	8	17	0	NA	NA						
	1,2,3-Trichlorobenzene	15	59	6	0.00017	0.00013	49					
	1,2,3-Trichloropropane	15	59	0	NA	NA	0.005		11			
	1,2,4-Trimethylbenzene	15	59	9	13.2	0.014	62					
	1,2-Dibromo-3-chloropropane	15	59	0	NA	NA	0.0054		18			
	1,2-Dibromoethane (EDB)	15	59	0	NA	NA	0.034		11			
	1,2-Dichloroethane (EDC)	15	59	0	NA	NA	0.43		4			
	1,2-Dichloropropane	15	58	0	NA	NA	0.94		2			
	1,3,5-Trimethylbenzene	15	59	8	5.5	0.026	780					
	1,3-Dichlorobenzene	7	40	0	NA	NA	37.7					
	1,3-Dichloropropane	8	17	0	NA	NA	1600					
	1,4-Difluorobenzene	1	1	1	2	2						
VOCs	2,2-Dichloropropane	8	17	0	NA	NA						
	2-Butanone	15	59	2	2.4	0.015	28000					
	2-Chlorotoluene	8	17	0	NA	NA	1600					
	2-Hexanone	15	59	0	NA	NA	12.6		2			
	4-Chlorotoluene	8	17	0	NA	NA	1600					
	4-Methyl-2-pentanone	15	59	0	NA	NA	5300					
	Acetone	15	59	30	0.064	0.0065	61000					
	Benzene	15	59	22	12	0.00069	1.1	3				
	Bromobenzene	8	17	0	NA	NA	300					
	Bromochloromethane	15	59	0	NA	NA	160					
	Bromodichloromethane	15	59	0	NA	NA	0.27		5			
	Bromoform	15	59	0	NA	NA	15.9					
	Bromomethane	15	58	0	NA	NA	7.3					
	Carbon disulfide	15	59	4	0.0075	0.0043	820					
	Carbon tetrachloride	15	59	0	NA	NA	0.61		2			
	Chlorobenzene	15	59	0	NA	NA	290					
F	Chlorobenzene-d5	1	1	1	2	2						
ļ	Chloroethane	15	59	0	NA	NA	15000					
F	Chloroform	15	59	3	0.044	0.00048	0.29		5			
ŀ	Chloromethane	15	59	0	NA	NA	120					

#### Table 7-3 - Statistical Summary of Soil Data

Bremerton Gas Works Site Bremerton, Washington

					Maximum	Minimum		Detected	Results with Reporting	Puget Sound	Number of Detected	Number of Non-Detect Results
1 1					Detected	Detected		Concentrations	Limit Concentrations	Background Metals	Concentrations	with Reporting Limit
Chemical		Number of	Number of	Number of			Soil Initial PRG	Exceeding the	that Exceed the Initial	Concentration	Exceeding Puget	Concentrations that Exceed
Group	Chemical Constituent	Locations	Samples	Detections	(mg/kg)	(mg/kg)	(mg/kg)	Initial PRG	Soil PRG	(mg/kg) <sup>1</sup>	Sound Background	Puget Sound Background
	cis-1,2-Dichloroethene (DCE)	15	59	0	NA	NA	160					
1 1	cis-1,3-Dichloropropene	15	59	3	0.93	0.00063	0.398	1	4			
1 1	Cyclohexane	7	42	0	NA	NA	7000					
1 1	Dibromochloromethane	15	59	0	NA	NA	0.68		2			
1 [	Dibromomethane	8	17	0	NA	NA	25					
	Dichlorodifluoromethane	15	59	0	NA	NA	94					
	Ethylbenzene	15	59	16	24	0.00073	5.4	1				
	Hexachlorobutadiene	1	1	0	NA	NA	6.2					
	Hexachloroethane	15	56	0	NA	NA	12		2			
1 [	Isopropylbenzene	15	59	7	1.6	0.00094	2100					
1 [	Methyl acetate	7	42	1	0.16	0.16	78000					
	Methyl tert-butyl ether (MTBE)	15	59	0	NA	NA	43					
	Methylcyclohexane	7	42	3	0.0038	0.00037						
	Methylene chloride	15	59	24	1.3	0.00058	56					
	n-Butylbenzene	8	17	2	1.96	1.78	3900					
VOCs	n-Hexane	8	17	1	0.00121	0.00121	570					
(continued)	n-Propylbenzene	8	17	2	0.952	0.792	3400					
	Pentafluorobenzene	2	3	3	2	0.04						
[	p-Isopropyltoluene	8	17	4	1.65	0.493						
[	sec-Butylbenzene	8	17	2	0.915	0.748	7800					
1 [	Styrene	15	59	4	0.07	0.000814	6300					
	tert-Butylbenzene	8	17	0	NA	NA	7800					
	Tetrachloroethene (PCE)	15	59	3	0.00059	0.00044	22					
1 1	Toluene	15	59	30	7.5	0.00026	5000					
	trans-1,2-Dichloroethene	15	59	0	NA	NA	150					
	trans-1,3-Dichloropropene	15	59	3	0.93	0.00063	0.398	1	4			
	Trichloroethene (TCE)	15	59	3	0.00147	0.00044	0.91		2			
	Trichlorofluoromethane	15	59	13	0.0078	0.0006	790					
	Vinyl chloride	15	59	0	NA	NA	0.06		11			
	m,p-Xylenes	13	50	9	57	0.00052	630					
	o-Xylene	13	50	8	55	0.00049	690					
	Xylenes (total)	8	17	7	16.7	0.353	630					
1 1	Aroclor 1016	8	17	0	NA	NA	3.9					
1 1	Aroclor 1221	8	17	0	NA	NA	0.14					
1 1	Aroclor 1232	8	17	0	NA	NA	0.14					
	Aroclor 1242	8	17	0	NA	NA	0.22					
PCBs	Aroclor 1248	8	17	0	NA	NA	0.22					
	Aroclor 1254	8	17	0	NA	NA	0.22					
	Aroclor 1260	8	17	0	NA	NA	0.22					
	Aroclor 1262	8	17	0	NA	NA						
	Aroclor 1268	8	17	0	NA	NA						

#### Notes:

<sup>1</sup> Background metals concentrations based on Puget Sound (when available) or Washington State background (Ecology 1994).

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

PAHs = polycyclic aromatic hydrocarbons

mg/kg = millograms per kilogram

PCBs = polychlorinated biphenyls

PRG = preliminary remediation goal

SVOCs = semi-volatile organic compounds

NA = Not applicable, as there are no detections.

TPH = total petroleum hydrocarbons VOCs = volatile organic compounds

Bremerton Gas Works Site Bremerton, Washington

					Maximum Detected	Minimum Detected	Groundwater	Number of Detected Concentrations Exceeding the	Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the	Surface Water Initial	Number of Detected Concentrations Exceeding the	Number of Non-Detect Results with Reporting Limit Concentrations that Exceed the
Chemical		Number of	Number of	Number of	Concentration	Concentration	Initial PRG	Groundwater Initial	Groundwater Initial	PRG	Surface Water	Surface Water Initial
Group	Chemical Constituent	Locations	Samples	Detections	(ug/L)	(ug/L)	(ug/L)	PRG	PRG	(ug/L)	Initial PRG	PRG
	Gasoline Range Hydrocarbons	10	10	7	10600	63.5						
TPH	Diesel Range Hydrocarbons	11	11	6	18500	170						
	Oil Range Hydrocarbons	11	11	1	160	160						
	Antimony	10	10	2	0.4	0.3	6			640		
	Arsenic	10	10	10	26	0.6	0.045	10		0.14	10	
	Barium	2	2	2	173	35.7	2000					
	Beryllium	10	10	3	1.08	0.37	4			0.66	2	7
	Cadmium	10	10	2	0.16	0.05	5			8.8		
	Chromium (Total)	10	10	10	228	1.34	100	2		42	3	
	Chromium (VI)	8	8	7	90	6	0.031	7	1	50	2	
	Cobalt	2	2	2	8.3	1.4	4.7	1				
Metals (Total)	Copper	10	10	10	143	1.05	620			3.1	8	
ivietais (Total)	Lead	10	10	8	21.6	0.44	15	2		8.1	2	
	Manganese	2	2	2	3020	98.1	320	1				
	Mercury	8	8	1	0.246	0.246	0.63			0.94		
	Nickel	10	10	10	232	1.65	300			8.2	7	
	Selenium	10	10	1	3.64	3.64	50			71		
	Silver	10	10	1	0.07	0.07	71			1.9		
	Thallium	10	10	1	0.26	0.26	0.16	1	9	0.47		9
	Vanadium	2	2	2	78.2	3.7	63	1				
	Zinc	10	10	8	185	4.5	4700			81	2	
	Acenaphthene	9	9	5	485	1.1	400	1		990		
	Acenaphthylene	10	10	6	34.9	0.222				4840		
	Anthracene	10	10	5	120	0.4	1300			40000		
	Benzo(g,h,i)perylene	10	10	5	25.6	0.0979				7.64	1	
	Dibenzofuran	10	10	2	31.8	0.29	5.8	1	7	4	1	7
PAHs	Fluoranthene	10	10	6	122	0.26	630			140		
174.5	Fluorene	10	10	7	184	0.102	220			5300		
	Phenanthrene	10	10	5	377	1.04				1.5	3	
	Pyrene	10	10	7	34.5	0.174	87			4000		
	1-Methylnaphthalene	8	8	4	970	0.813	0.97	3	1	2.1	3	1
	2-Methylnaphthalene	10	10	6	1430	0.13	27	1		4.2	1	
	Naphthalene	2	2	0	NA	NA	0.14			13		
	Benz(a) anthracene	10	10	6	39.3	0.0168	0.029	5	2	0.018	5	2
	Benzo(a) pyrene	10	10	6	37.6	0.0247	0.0029	6	4	0.018	6	2
	Benzo(b)fluoranthene	10	10	4	0.657	0.0968	0.029	4	3	0.018	4	3
	Benzo(k)fluoranthene	10	10	5	0.615	0.0602	0.29	2	1	0.018	5	3
cPAHs	Chrysene	10	10	6	40.8	0.0372	2.9	1		0.018	6	2
	Dibenzo(a,h)anthracene	10	10	4	0.189	0.0437	0.0029	4	6	0.018	4	3
	Indeno(1,2,3-cd)pyrene	10	10	4	0.467	0.0874	0.029	4	3	0.018	4	3
	Total cPAHs TEQ (ND = 0)	10	10	6	41.9	0.0328	0.0029	6				
	Total cPAHs TEQ (ND = 1/2 RDL)	10	10	6	43.8	0.0342	0.0029	6				

Bremerton Gas Works Site Bremerton, Washington

Chemical	Chamia l Canatita and	Number of Locations	1 1	Number of	Maximum Detected Concentration	Minimum Detected Concentration	Groundwater Initial PRG	Number of Detected Concentrations Exceeding the Groundwater Initial PRG	Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the Groundwater Initial PRG	Surface Water Initial PRG	Number of Detected Concentrations Exceeding the Surface Water Initial PRG	Number of Non-Detect Results with Reporting Limit Concentrations that Exceed the Surface Water Initial PRG
Group	Chemical Constituent		Samples	Detections	(ug/L)	(ug/L)	(ug/L)	PNG	PNG	(ug/L)	IIIIIIIII PNG	PNG
	1,1'-Biphenyl	2	2	0	NA NA	NA NA	0.83 1.2			14 1.1		
	1,2,4,5-Tetrachlorobenzene 1,2,4-Trichlorobenzene	2 10	2 10	0	NA NA	NA NA	0.99		8	70		
Other SVOCs	1,2,4-Trimethylbenzene	1	1	0	NA NA	NA NA	15			19		
	1,2-Dichlorobenzene	10	10	0	NA	NA	280			1300		
	1,3-Dichlorobenzene	10	10	0	NA	NA				960		
	1,4-Dichlorobenzene	10	10	0	NA	NA	0.42			190		
	2,3,4,6-Tetrachlorophenol	2	2	0	NA	NA	170			1.2		
	2,4,5-Trichlorophenol	10	10	0	NA 	NA	890		_	12		
	2,4,6-Trichlorophenol 2,4-Dichlorophenol	10 10	10 10	0	NA NA	NA NA	3.5 35		8	2.4 290		8
	2,4-Dimethylphenol	10	10	0	NA NA	NA NA	270			850		
	2,4-Dinitrophenol	10	10	0	NA NA	NA NA	30			5300		
	2-Chloronaphthalene	10	10	0	NA NA	NA NA	550			1600		
	2-Chlorophenol	10	10	0	NA	NA	71			150		
	2-Methylphenol	8	8	0	NA	NA	720			67		
	2-Nitroaniline	10	10	0	NA	NA	150					
	2-Nitrophenol	10	10	0	NA	NA 				2940		
	3 & 4 Methylphenol 3,3'-Dichlorobenzidine	8 10	8 10	0	NA NA	NA NA	1400 0.11	-	10	0.028		10
	3,3 -Dichlorobenzidine 3-Nitroaniline	10	10	0	NA NA	NA NA	0.11		10	0.028		10
	4,6-Dinitro-2-methylphenol	10	10	0	NA NA	NA NA	1.2		8	280		
	4-Bromophenyl phenyl ether	10	10	0	NA	NA			<u> </u>	1.5		8
	4-Chloro-3-methylphenol	10	10	0	NA	NA	1100			34.8		
	4-Chloroaniline	10	10	0	NA	NA	0.32		10	232		
	4-Chlorophenyl phenyl ether	10	10	0	NA	NA						
	4-Methylphenol	2	2	0	NA	NA	1400			25		
	4-Nitroaniline 4-Nitrophenol	10 10	10 10	0	NA NA	NA NA	3.3		8	60		
	4-Nitrophenoi Acenaphthene	9	9	5	NA 485	1.1	400	1		990		
	Acetophenone	2	2	0	NA	NA NA	1500	-		330		
	Aniline	8	8	0	NA	NA	12			2.2		8
	Atrazine	2	2	0	NA	NA	0.26		2	1.8		
Other SVOCs	Benzaldehyde	2	2	0	NA	NA	1500					
(continued)	Benzidine	2	2	0	NA	NA	0.000092		2	3.9		
	Benzoic acid	8	8	0	NA	NA	58000			42		_
	Benzyl alcohol	8 10	8 10	0	NA 0.33	NA 0.33	1500 14			8.6 1900		8
	Benzyl butyl phthalate Bis(2-chloro-1-methylethyl) ether	10	10	0	0.33 NA	0.33 NA	0.31		10	65000		
	Bis(2-chloroethoxy)methane	10	10	0	NA NA	NA NA	46		10	03000		
	Bis(2-chloroethyl) ether	10	10	0	NA NA	NA NA	0.012		10	0.53		8
	Bis(2-ethylhexyl) phthalate	10	10	2	0.5	0.33	4.8		8	2.2		8
	Caprolactam	2	2	1	0.71	0.71	7700					
	Carbazole	10	10	1	1.3	1.3						
	Diethyl phthalate	10	10	0	NA 	NA 	11000			44000		
	Dimethyl phthalate	10	10	0	NA NA	NA NA	670			1100000		
	Di-n-butyl phthalate Di-n-octyl phthalate	10 10	10 10	0	NA NA	NA NA	670 160			4500 22		
	Hexachlorobenzene	10	10	0	NA NA	NA NA	0.042		10	0.00029		10
	Hexachlorobutadiene	10	10	0	NA NA	NA NA	0.26		8	18		10
	Hexachlorocyclopentadiene	10	10	0	NA NA	NA NA	22			1100		
	Hexachloroethane	10	10	0	NA	NA	0.79		8	3.3		8
	Isophorone	10	10	0	NA	NA	67			960		
	Nitrobenzene	8	8	0	NA	NA	0.12		8	690		
	N-Nitrosodimethylamine	2	2	0	NA 	NA 	0.00042		2	330000		
	N-Nitroso-di-n-propylamine	10	10	0	NA NA	NA NA	0.0093		10	0.51		8
	N-Nitrosodiphenylamine  Pentachlorophenol	10 10	10 10	2	NA 11.4	NA 0.1	10 0.035	2	1 8	6 3	1	8 7
	Phenol	10	10	3	81.6	75.5	4500		0	860000	1	<del>'</del>
	2,4-Dinitrotoluene	8	8	0	NA NA	NA NA	0.2		8	3.4		8
	2,6-Dinitrotoluene	8	8	0	NA NA	NA NA	0.042		8	81		
	2-Methylnaphthalene	10	10	6	1430	0.13	27	1		4.2	1	

Bremerton Gas Works Site Bremerton, Washington

Chemical Group	Chemical Constituent	Number of Locations	Number of Samples	Number of Detections	Maximum Detected Concentration (ug/L)	Minimum Detected Concentration (ug/L)	Groundwater Initial PRG (ug/L)	Number of Detected Concentrations Exceeding the Groundwater Initial PRG	Number of Non- Detect Results with Reporting Limit Concentrations that Exceed the Groundwater Initial PRG	Surface Water Initial PRG (ug/L)	Number of Detected Concentrations Exceeding the Surface Water Initial PRG	Number of Non-Detect Results with Reporting Limit Concentrations that Exceed the Surface Water Initial PRG
Стоир								i riko	FING	(46/1/	IIIIdai Filo	FING
	1,1,1,2-Tetrachloroethane	10	10	0	NA NA	NA NA	0.5			70		
	1,1,1-Trichloroethane	10	10	0	NA NA	NA NA	200			76		
	1,1,2 - Trichlorotrifluoroethane 1,1,2,2-Tetrachloroethane	10	2 10	0	NA NA	NA NA	53000 0.066		10	4		
	1,1,2,7-retraction oethane	10	10	0	NA NA	NA NA	0.24		2	16		
	1,1-Dichloroethane	10	10	0	NA NA	NA NA	2.4		2	47		
	1,1-Dichloroethene	9	9	0	NA NA	NA NA	7			7100		
	1,1-Dichloropropene	8	8	0	NA NA	NA NA	,			7100		
	1,2,3-Trichlorobenzene	10	10	0	NA NA	NA NA	5.2			8		
	1,2,3-Trichloropropane	10	10	0	NA NA	NA NA	0.00065		10	-		
	1,2,4-Trichlorobenzene	10	10	0	NA	NA	0.99		8	70		
	1,2,4-Trimethylbenzene	1	1	0	NA	NA	15			19		
	1,2-Dibromo-3-chloropropane	10	10	0	NA	NA	0.00032		10			
	1,2-Dibromoethane (EDB)	10	10	0	NA	NA	0.0065		10			
	1,2-Dichlorobenzene	10	10	0	NA	NA	280			1300		
	1,2-Dichloroethane (EDC)	10	10	3	4.72	0.93	0.15	3	7	37		
	1,2-Dichloropropane	10	10	0	NA	NA	0.38			15		
	1,3,5-Trimethylbenzene	10	10	5	30	0.53	87			71		
	1,3-Dichlorobenzene	10	10	0	NA	NA				960		
	1,3-Dichloropropane	8	8	0	NA	NA	290					
	1,4-Dichlorobenzene	10	10	0	NA	NA	0.42			190		
	2,2-Dichloropropane	8	8	0	NA	NA						
	2-Butanone	10	10	0	NA	NA	4900			2200		
	2-Chlorotoluene	8	8	0	NA	NA	180					
	2-Hexanone	10	10	0	NA	NA	34			99		
	4-Chlorotoluene	8	8	0	NA	NA	190					
	4-Methyl-2-pentanone	10	10	0	NA	NA	1000			170		
VOCs	Acetone	10	10	0	NA 	NA	12000	_		1700	_	
	Benzene	10	10	8	950	2.23	0.39	8		51	5	
	Bromobenzene	8	8	0	NA NA	NA NA	54 83			-		
	Bromochloromethane Bromodichloromethane	10 10	10 10	0	NA NA	NA NA	0.12		10	17		
	Bromoform	10	10	0	NA NA	NA NA	7.9		10	140		
	Bromomethane	10	10	0	NA NA	NA NA	7.9			1500		
	Carbon disulfide	10	10	0	NA NA	NA NA	720			0.92		
	Carbon disunde  Carbon tetrachloride	10	10	1	0.66	0.66	0.39	1		1.6		
	Chlorobenzene	10	10	0	NA	NA	72	<del>                                     </del>		1600		
	Chloroethane	10	10	0	NA NA	NA NA	21000					
	Chloroform	10	10	3	2.84	0.2	0.19	3	7	470		
	Chloromethane	10	10	0	NA NA	NA	190	-		2700		
	cis-1,2-Dichloroethene (DCE)	10	10	3	1.29	0.37	28					
	cis-1,3-Dichloropropene	10	10	0	NA	NA				21		
	Cyclohexane	2	2	1	0.38	0.38	13000					
	Dibromochloromethane	10	10	0	NA	NA	0.15		10	13		
	Dibromomethane	8	8	0	NA	NA	7.9					
	Dichlorodifluoromethane	10	10	0	NA	NA	190					
	Ethylbenzene	10	10	7	322	0.53	1.3	6		2100		
	Hexachlorobutadiene	10	10	0	NA	NA	0.26		8	18		
	Hexachloroethane	10	10	0	NA	NA	0.79		8	3.3		8
	Isopropylbenzene	10	10	6	37.4	3	390			2.6	6	
	Methyl acetate	2	2	0	NA	NA	16000					
	Methyl tert-butyl ether (MTBE)	10	10	0	NA	NA	12			11070		
	Methylcyclohexane	2	2	0	NA	NA						
	Methylene chloride	10	10	0	NA	NA	5			590		
	n-Butylbenzene	8	8	4	5.3	0.48	780					_
	n-Hexane	8	8	1	1.17	1.17	250			0.58	1	7

Bremerton Gas Works Site Bremerton, Washington

									Number of Non-			
									Detect Results with		Number of	Number of Non-Detect
								Number of Detected	Reporting Limit		Detected	Results with Reporting
					Maximum	Minimum		Concentrations	Concentrations that	Surface	Concentrations	Limit Concentrations
					Detected	Detected	Groundwater	Exceeding the	Exceed the	Water Initial	Exceeding the	that Exceed the
Chemical		Number of	Number of	Number of	Concentration	Concentration	Initial PRG	Groundwater Initial	Groundwater Initial	PRG	Surface Water	Surface Water Initial
Group	Chemical Constituent	Locations	Samples	Detections	(ug/L)	(ug/L)	(ug/L)	PRG	PRG	(ug/L)	Initial PRG	PRG
Стопр		_						FNG	FNG		IIIIIIIII FRG	PNG
	n-Propylbenzene	8	8	4	9.2	2.38	530			128		
	p-Isopropyltoluene	8	8	4	8.44	0.27				85		
	sec-Butylbenzene	8	8	5	4.43	0.32	1600					
	Styrene	10	10	0	NA	NA	100			32		
	tert-Butylbenzene	8	8	0	NA	NA	510					
	Tetrachloroethene (PCE)	10	10	0	NA	NA	5			3.3		
VOCs	Toluene	10	10	6	41.9	0.45	860			15000		
(continued)	trans-1,2-Dichloroethene	10	10	0	NA	NA	86			10000		
(,	trans-1,3-Dichloropropene	10	10	0	NA	NA				21		
	Trichloroethene (TCE)	10	10	6	4.79	0.33	0.44	4		30		
	Trichlorofluoromethane	10	10	0	NA	NA	1100					
	Vinyl chloride	10	10	0	NA	NA	0.015		10	2.4		
	m,p-Xylenes	10	10	6	383	0.74	190	1				
	o-Xylene	10	10	6	211	4.91	190	1				
	Xylenes (total)	8	8	5	593	8.29	190	2		19	4	
	Aroclor 1016	8	8	0	NA	NA	0.96					
	Aroclor 1221	8	8	0	NA	NA	0.004		8			
	Aroclor 1232	8	8	0	NA	NA	0.004		8			
	Aroclor 1242	8	8	0	NA	NA	0.034		8			
PCBs	Aroclor 1248	8	8	0	NA	NA	0.034		8			
	Aroclor 1254	8	8	0	NA	NA	0.034		8			
	Aroclor 1260	8	8	0	NA	NA	0.034		8			
	Aroclor 1262	8	8	0	NA	NA				•		
	Aroclor 1268	8	8	0	NA	NA						

#### Notes:

cPAHs = carcinogenic polycyclic aromatic hydrocarbons NA = Not applicable, as there are no detections. PAHs = polycyclic aromatic hydrocarbons PCBs = polychlorinated biphenyls PRG = preliminary remediation goal SVOCs = semi-volatile organic compounds TPH = total petroleum hydrocarbons ug/L = micrograms per liter

VOCs = volatile organic compounds

### **Table 7-5 - Statistical Summary of Sediment Data** Bremerton Gas Works Site Bremerton, WA

Chemical Group	Chemical Constituent	Number of Locations	Number of Samples	Number of Detections	Maximum Detected Concentration (ug/kg)	Minimum Detected Concentration (ug/kg)	Sediment Initial PRG (ug/kg)	Puget Sound Background Sediment Metals Concentration <sup>1</sup> (ug/kg)	Number of Detected Concentrations Exceeding the Sediment Initial PRG	Number of Detected Concentrations Exceeding Puget Sound Background Metals Concentration
	Gasoline Range Hydrocarbons	5	5	0	NA NA	NA NA	(**8,7*8,7 	(**8,**8)		
TPH	Diesel Range Hydrocarbons	5	5	4	240000	63000				
	Oil Range Hydrocarbons	5	5	5	620000	21000				
	Aluminum	5	5	5	9030000	6020000				
	Antimony	1	1	1	3900	3900	2000	5000	1	
	Arsenic	5	5	5	5100	1500	57000	11000	-	
	Barium	5	5	5	47000	13300	37000	11000		
	Beryllium	5	5	5	2700	1900				
	Cadmium	5	5	0	NA NA	NA NA	5100	1000		
	Calcium	5	5	5	33600000	2390000	3100	1000		
	Chromium (Total)	5	5	5	21200	16600	260000	62000		
	Cobalt	5	5	5	26300	3000	50000	11000		
	Copper	5	5	5	71700	8600	390000	44000		
	Iron	5	5	5	15900000	9730000	20000000	44000		
Metals	Lead	5	5	5	30000	8900	450000	21000		
	Magnesium	5	5	5	4640000	3350000	450000	21000		
	Manganese	5	5	5	180000	135000	460000			
	Mercury	3	3	3	100	27.8	410	200		
	Nickel	5	5	5	52600	21400	20900	50000	5	1
	Potassium	5	5	5	603000	415000	20300	30000		<u> </u>
	Selenium	5	5	1	400	400	2000	780		
	Silver	5	5	0	NA NA	NA NA	6100	300		
	Sodium	5	5	5	1930000	605000	0100	300		
	Thallium	5	5	0	NA NA	NA NA				
	Vanadium	5	5	5	36500	21600		45000		
	Zinc	5	5	5	79900	23200	410000	93000		
	Acenaphthene	48	63	61	160000	0.4	500	93000	16	
	Acenaphthylene	51	66	66	840000	0.7	1300		33	
	Anthracene	51	66	66	680000	0.3	960		41	
	Benzo(g,h,i)perylene	51	66	66	260000	0.9	670		50	
	Dibenzofuran	5	5	4	74	58	540		30	
	Fluoranthene	46	61	61	1100000	1.6	1700		45	
	Fluorene	51	66	65	600000	0.3	540		36	
	Phenanthrene	51	66	66	1700000	2.6	1500		46	
	Pyrene	51	66	66	1400000	1.6	2600		48	
	2-Methylnaphthalene	5	5	5	1200	19	670		1	
	Naphthalene	46	61	61	1700000	5.4	2100		23	
PAHs	Benz(a)anthracene	51	66	66	310000	0.3	1300		46	
17413	Benz(a)antriracene Benzo(a)pyrene	51	66	66	400000	0.5	1600		47	
	Benzo(b)fluoranthene	51	66	66	200000	0.3	10400		17	<del> </del>
	Benzo(b)fluoranthene Benzo(k)fluoranthene	51	66	65	93000	0.4	240		50	1
	Chrysene	51	66	66	270000	0.5	1400		47	<del> </del>
	Dibenzo(a,h)anthracene	51	66	65	38000	0.3	230		46	<del> </del>
	Indeno(1,2,3-cd)pyrene	51	66	66	190000	0.2	600		49	
	Total cPAHs TEQ (ND = 0)	51	66	66	509200	0.4	1600		49	<del> </del>
	Total cPAHs TEQ (ND = 1/2 RDL)	51	66	66	509200	0.9	1600		49	
	Total HPAHs  Total HPAHs	46	61	61	4361000	6.2	12000		45	<del> </del>
	Total LPAHs	46	61	61	5596000	10.1	5200		39	
	Total PAHs	46	61	61	8890000	16.3	4022		48	-

### **Table 7-5 - Statistical Summary of Sediment Data**Bremerton Gas Works Site Bremerton, WA

Chemical Group	Chemical Constituent	Number of Locations	Number of Samples	Number of Detections	Maximum Detected Concentration (ug/kg)	Minimum Detected Concentration (ug/kg)	Sediment Initial PRG (ug/kg)	Puget Sound Background Sediment Metals Concentration <sup>1</sup> (ug/kg)	Number of Detected Concentrations Exceeding the Sediment Initial PRG	Number of Detected Concentrations Exceeding Puget Sound Background Metals Concentration
	1,1'-Biphenyl	5	5	4	110	60	1220	1		
	1,2,4,5-Tetrachlorobenzene	5	5	0	NA NA	NA NA	47000			
	1,2,4-Trichlorobenzene	8	9	0	NA NA	NA NA	31			
	1,2-Dichlorobenzene	8	9	0	NA NA	NA NA	35			
	1,3,5-Trimethylbenzene	1	1	1	21	21	- 33			
	1,3-Dichlorobenzene	5	5	0	NA NA	NA NA	842			
	1,4-Dichlorobenzene	2	2	2	23	22	110			
	1,4-Dioxane	5	5	0	NA NA	NA NA	119			
	2,3,4,6-Tetrachlorophenol	5	5	0	NA NA	NA NA	284			
	2,4,5-Trichlorophenol	5	5	0	NA NA	NA NA	819			
	2,4,6-Trichlorophenol	5	5	0	NA NA	NA NA	2650			
	2,4-Dichlorophenol	5	5	0	NA NA	NA NA	117			
	2,4-Dimethylphenol	5	5	0	NA NA	NA NA	29			
	2,4-Dinitrophenol	5	5	0	NA NA	NA NA	6.21			
	2-Chloronaphthalene	5	5	0	NA NA	NA NA	417			
	2-Chlorophenol	5	5	0	NA NA	NA NA	344			
	2-Chlorophenol 2-Nitroaniline	5	5	0	NA NA	NA NA	344			
				0	1					
	2-Nitrophenol	5	5		NA NA	NA NA	2000			
	3,3'-Dichlorobenzidine	5	5	0	NA NA	NA NA	2060			
	3-Nitroaniline	5	5	0	NA	NA NA	404			
	4,6-Dinitro-2-methylphenol	5	5	0	NA	NA	104			
	4-Bromophenyl phenyl ether	5	5	0	NA	NA NA	1230			
	4-Chloro-3-methylphenol	5	5	0	NA	NA	388			
Oil	4-Chloroaniline	5	5	0	NA	NA	146			
Other	4-Chlorophenyl phenyl ether	5	5	0	NA	NA 1 =				
SVOCs	4-Methylphenol	5	5	2	17	17	670			
	4-Nitroaniline	5	5	0	NA	NA				
	4-Nitrophenol	5	5	0	NA	NA	13.3			
	Acenaphthene	48	63	61	160000	0.4	500		16	
	Acetophenone	5	5	0	NA	NA				
	Atrazine	5	5	0	NA	NA	6.62			
	Benzaldehyde	5	5	2	38	19				
	Benzidine	5	5	0	NA	NA				
	Benzyl butyl phthalate	5	5	0	NA	NA	63			
	Bis(2-chloro-1-methylethyl) ethe	5	5	0	NA	NA	ļ			
	Bis (2-chloroethoxy) methane	5	5	0	NA	NA				
	Bis(2-chloroethyl) ether	5	5	0	NA	NA	3520			
	Bis(2-ethylhexyl) phthalate	5	5	1	42	42	1300			
	Caprolactam	5	5	0	NA	NA				
	Carbazole	5	5	4	110	69	ļ			
	Dibenzofuran	5	5	4	74	58	540			
	Diethyl phthalate	5	5	0	NA	NA	200			
	Dimethyl phthalate	5	5	0	NA	NA	71			
	Di-n-butyl phthalate	5	5	0	NA	NA	1400			
	Di-n-octyl phthalate	5	5	0	NA	NA	6200			
	Hexachlorobenzene	5	5	0	NA	NA	22			
	Hexachlorobutadiene	8	9	0	NA	NA	11			
	Hexachlorocyclopentadiene	5	5	0	NA	NA	139			
	Hexachloroethane	3	3	0	NA	NA	804			
	Isophorone	5	5	0	NA	NA	432			

### **Table 7-5 - Statistical Summary of Sediment Data**Bremerton Gas Works Site Bremerton, WA

Chemical Group	Chemical Constituent	Number of Locations	Number of Samples	Number of Detections	Maximum Detected Concentration (ug/kg)	Minimum Detected Concentration (ug/kg)	Sediment Initial PRG (ug/kg)	Puget Sound Background Sediment Metals Concentration <sup>1</sup> (ug/kg)	Number of Detected Concentrations Exceeding the Sediment Initial PRG	Number of Detected Concentrations Exceeding Puget Sound Background Metals Concentration
	Naphthalene	46	61	61	1700000	5.4	2100		23	
	N-Nitrosodimethylamine	5	5	0	NA	NA				
Other	N-Nitroso-di-n-propylamine	5	5	0	NA	NA				
SVOCs	N-Nitrosodiphenylamine	5	5	0	NA	NA	28			
(continued)	Pentachlorophenol	5	5	5	110	35	360			
	Phenol	5	5	0	NA	NA	420			
	2-Methylnaphthalene	5	5	5	1200	19	670		1	
	1,1,1,2-Tetrachloroethane	8	9	0	NA	NA				
	1,1,1-Trichloroethane	8	9	0	NA	NA	856			
	1,1,2 - Trichlorotrifluoroethane	8	9	0	NA	NA				
	1,1,2,2-Tetrachloroethane	8	9	0	NA	NA	202			
	1,1,2-Trichloroethane	8	9	0	NA	NA	570			
	1,1-Dichloroethane	8	9	0	NA	NA	0.575			
	1,1-Dichloroethene	8	9	0	NA	NA	2780			
	1,1-Dichloropropene	3	4	0	NA	NA				
	1,2,3-Trichlorobenzene	8	9	0	NA	NA	858			
	1,2,3-Trichloropropane	8	9	0	NA	NA				
	1,2,4-Trichlorobenzene	8	9	0	NA	NA	31			
	1,2,4-Trimethylbenzene	8	9	4	980	2.4				
	1,2-Dibromo-3-chloropropane	8	9	0	NA	NA				
	1,2-Dibromoethane (EDB)	8	9	0	NA	NA				
	1,2-Dichlorobenzene	8	9	0	NA	NA	35			
	1,2-Dichloroethane (EDC)	8	9	0	NA	NA	260			
	1,2-Dichloropropane	8	9	0	NA	NA	333			
	1,3,5-Trimethylbenzene	1	1	1	21	21				
	1,3-Dichlorobenzene	5	5	0	NA	NA	842			
	1,3-Dichloropropane	3	4	0	NA	NA				
	1,4-Dichloro-2-Butene	3	4	0	NA	NA				
VOCs	1,4-Dichlorobenzene	2	2	2	23	22	110			
	2,2-Dichloropropane	3	4	0	NA	NA				
	2-Butanone	8	9	0	NA	NA	42.4			
	2-Chloroethyl Vinyl Ether	3	4	0	NA	NA				
	2-Chlorotoluene	3	4	0	NA	NA				
	2-Hexanone	8	9	0	NA	NA	58.2			
	4-Chlorotoluene	3	4	0	NA	NA				
	4-Methyl-2-pentanone	8	9	0	NA	NA	25.1			
	Acrolein	3	4	0	NA	NA	0.00152			
	Acrylonitrile	3	4	0	NA	NA	1.2			
	Benzene	8	9	3	8.1	1.5	137			
	Bromobenzene	3	4	0	NA	NA NA				
	Bromochloromethane	8	9	0	NA	NA NA	1			
	Bromodichloromethane	8	9	0	NA NA	NA NA	1		1	
	Bromoethane	3	4	0	NA NA	NA NA	1			
	Bromoform	8	9	0	NA NA	NA NA	1310			
	Bromomethane	8	9	0	NA NA	NA NA	1.37		1	
	Carbon disulfide	8	9	1	4.3	4.3	0.851		1*	
	Carbon tetrachloride	8	9	0	NA	NA	7240		<u> </u>	
	Chlorobenzene	 8	9	0	NA NA	NA NA	162			
	Chloroethane	 8	9	0	NA NA	NA NA	102			
	Chloroform	<u> </u>	9	0	NA NA	NA NA	121			

#### Table 7-5 - Statistical Summary of Sediment Data

Bremerton Gas Works Site Bremerton, WA

Chemical Group	Chemical Constituent	Number of Locations	Number of Samples	Number of Detections	Maximum Detected Concentration (ug/kg)	Minimum Detected Concentration (ug/kg)	Sediment Initial PRG (ug/kg)	Puget Sound Background Sediment Metals Concentration <sup>1</sup> (ug/kg)	Number of Detected Concentrations Exceeding the Sediment Initial PRG	Number of Detected Concentrations Exceeding Puget Sound Background Metals Concentration
	Chloromethane	8	9	0	NA	NA				
	cis-1,2-Dichloroethene (DCE)	8	9	0	NA	NA				
	cis-1,3-Dichloropropene	8	9	0	NA	NA				
	Cyclohexane	5	5	0	NA	NA				
	Dibromochloromethane	8	9	0	NA	NA				
	Dibromomethane	3	4	0	NA	NA				
	Dichlorodifluoromethane	5	5	0	NA	NA				
	Ethylbenzene	8	9	2	24	2.3	305			
	Hexachlorobutadiene	8	9	0	NA	NA	11			
	Hexachloroethane	3	3	0	NA	NA	804			
	Isopropylbenzene	8	9	2	9	0.48	86			
	Methyl acetate	5	5	0	NA	NA				
	Methyl tert-butyl ether (MTBE)	5	5	0	NA	NA				
	Methylcyclohexane	5	5	1	0.65	0.65				
	Methylene chloride	8	9	1	1.8	1.8	159			
VOCs	Methyliodide	3	4	0	NA	NA				
(continued)	n-Butylbenzene	3	4	1	84	84				
(continued)	n-Propylbenzene	3	4	1	8.3	8.3				
	p-Isopropyltoluene	3	4	0	NA	NA				
	sec-Butylbenzene	3	4	0	NA	NA				
	Styrene	8	9	0	NA	NA	7070			
	tert-Butylbenzene	3	4	0	NA	NA				
	Tetrachloroethene (PCE)	8	9	0	NA	NA	190			
	Toluene	8	9	2	1.5	0.51	1090			
	trans-1,2-Dichloroethene	8	9	0	NA	NA	1050			
	trans-1,3-Dichloropropene	8	9	0	NA	NA				
	Trichloroethene (TCE)	8	9	0	NA	NA	8950			
	Trichlorofluoromethane	8	9	0	NA	NA				
	Vinyl acetate	3	4	0	NA	NA	13			
	Vinyl chloride	8	9	0	NA	NA	202			
	m,p-Xylenes	8	9	2	2.9	1.7				
	o-Xylene	8	9	2	5.7	3.9				
	Naphthalene	46	61	61	1700000	5.4	2100		23	

#### Notes:

\*Carbon disulfide is a common laboratory chemical. Based on the review of existing analytical data quality, these detections are considered to be the result of laboratory cross-contamination. The results are not considered representative of site conditions.

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

HPAH = high molecular weight PAH

LPAH = low molecular weight PAH

NA = Not applicable, as there are no detections.

PAHs = polycyclic aromatic hydrocarbons

PCBs = polychlorinated biphenyls

PRG = Preliminary Remediation Goal

SVOCs = semivolatile organic compounds

TPH = total petroleum hydrocarbons

VOCs = volatile organic compounds

ug/kg = micrograms per kilogram

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<sup>&</sup>lt;sup>1</sup> Background metals concentrations based on Puget Sound (when available) or Washington State background (Ecology 1994).

Bremeton Gas Works Site Bremerton, Washington

MGP Site Name & Location	Reference	Geologic Conditions	Groundwater / Surfacewater	Contaminants of Concern	Remedial Actions	Cleanup Status
Cold Spring MGP Ste Cold Spring, NY	Record of Decition (2010) http://www.dec.ny.gov/docs/r emediation_hudson_pdf/e34 0026arod.pdf	Subsurface soils consist of 11-13 feet of debris containing fill underlain by a 15 foot thick layer of clay, which overfles bedrock.  Contamination confined to the fill material.	*Groundwater flows to the west, towards the Hudson River which is adjacent to the site.  *No contamination was observed in river sediments.	BTEX PAHs	Excavation and off-site treatment/disposal.	Scheduled to begin late 2014
Saranac Street MGP Site Plattsburgh, NY	emediation_hudson_pdf/rod 51000701.pdf	Subsurface soils consist of up to 21 feet of debris containing fill underlain by up to 15 foot thick layer of sandy alluvium. Beneith the alluvium lies a layer of dense glacial till, which overlies limestone bedrock. Contamination present down to and into fractured bedrock.	*The Saranac River forms the southern, western, and northern site boundary.  *Coal tar discharged into the river along the northwestern and norther site boundaries.	BTEX PAHs	In situ stabilization; Soil and sediment excavation with off-site treatment/disposal; Bedrock tar collection wells.	Remedial Action complete
Waterville MFG Plant Waterville, NY	http://www.dec.ny.gov/docs/r emediation hudson pdf/6330 41_1.pdf	Subsurface soils consist of one foot of topsoil over a fill unit up to 12 feet thick consisting of a substantial amount of ash as well as brown sand and gravel, coal fragments and bricks. Below the fill is a unit of glacial outwash sand and silt ranging in thickness from 1 to 10 feet. A dense kame moraine silt and gravel deposit of depths from 4 to 12 feet was found below the outwash unit.  *Contamination present up to 14 feet below grade.	A western flowing tributary to Big Creek forms the southern edge of the property, approximately 150 feet south of the site.  The depth to groundwater ranges from approximately 4 to 12 feet below grade. Groundwater flow through the site is to the south-southwest and discharges into the Big Creek tributary.	BTEX PAHs	Excavation and Disposal; Institutional Controls; Soil Cap.	No Further Action required
Cortiand Homer Former MGP Site Homer, NY	amediation budges pdf/red7	Subsurface soils consist of a fill layer ranging from 6 inches to 10 feet and is underlain by outwash sand that varies in thickness from 20 to 40 feet. A confining silt/clay layer was observed benieth the outwash sand. Contamination present up to 37 feet below grade.	The West Branch of the Tioughnioga River is located 150 feet east of the site parcels.  Depth to groundwater at the site is approximately 5 feet below grade. Groundwater flow is in a east to east-southeast direction. Groundwater discharges into the river.  River sediments have been impacted by contaminants.	BTEX PAHs Cyanide	Excavation and disposal of source area soils;  In situ stabilization of downgradient contaminated soils;  NAPL collection trench;  Sediment removal.	Remedial Design complete

Bremeton Gas Works Site Bremerton, Washington

MGP Site Name & Location	Reference	Geologic Conditions	Groundwater / Surfacewater	Contaminants of Concern	Remedial Actions	Cleanup Status
Tacoma Tar Pits Tacoma, WA	http://yosemite.epa.gov/R10/ CLEANUP.NSF/sites/TacomaTar pis/SFLE/TTP-SYr-Review- Sept03.pdf	Subsurface soils consist of several feet of fill underlain by a layered sequence of silts and sands.	The Puyallup River is just norheast of the site. Groundwater occurs several feet below ground surface at the Tacoma Tar Pits site. The groundwater levels at the site vary in response to the tidal action in Commencement Bay and adjacent waterways. Groundwater flow directions vary depending on location, season, and tide stage. In general however, groundwater typically flows east (northwest and central potions of the site) and south (southeast portion of the site).	BTEX PAHs	Excavation and stabilization; Stabilized material placed in an engineered waste pile on site; Soil cap; Groundwater pump and treat.	Ongoing O&M for cover and groundwater treatment system
Oakland MGP Oakland, CA		Subsurface soils consisting of up to 5 feet of gravel/sand fill underlain by a sandy layer that extends up to 15 feet below grade with interbeded layers of silt and clay. The sandy layer is underlain by a fine-grained layer of clay and silt up to 20 feet below grade.  Contamination present up to 21 feet below grade.	Groundwater is 2 to 7:5 feet bgs and flows towards the Oakland Inner Harbor, which is approximately 1000 feet away.	TPH BTEX PAHs Cyanide	Soil cap.	Ongoing O&M
Glens Falls - Mohican Street MGP Glens Falls, NY	http://www.dec.ny.gov/docs/r emediation hudson pdf/5570 16roda2.pdf	Subsurface soil cosists of fill underlain by glacial fluvial deposits of sand, silt, silty sand, sandy silt. A layer of silty day overlies bedrock, which is encountered between 9-29 feet below grade. Contamination present up to 19 feet below grade.	The site is bounded to the south by the Glens Falls feeder canal.  Groundwater is 2-14 feet below grad and flows towards the Glens Falls canal and Hudson River.  Canal sediments are impacted.	BTEX PAHs	Excavation of source material; Oxygen delivery system; Soil cover; Institutional controls; Dredging and disposal.	Remedial Action approved
Gastown MGP Site Tonawanda, NY	http://www.dec.ny.gov/docs/r emediation hudson pdf/rod9 1517ttext.pdf http://www.dec.ny.gov/chemi cal/58387.html	Subsurface soils consist of up to 22 feet of debris containing fill undertaini by layers of sand and silt for an additional 24 feet below grade.  Contamination present down into the sand/silt layers.	The site is bounded to the north-northwest by Tonawanda Creek.  Groundwater is approximately 6 feet below grade and flows to the north into Tonawanda Creek.  Creek sediments have been impacted.	BTEX PAHs	Excavation and disposal;  In situ stabilization;  NAPL collection wells.	Scheduled to begin in 2013
Former Sacramento MGP Sacramento, CA	http://www.pge.com/about/e nvironment/taking- responsibility/mgp/sacramen to.shtml	Subsurface soils consist of up to 15 feet of fill undertain by a layer containing mostly silts and dayey silts to 25 feet below grade. A layer of unconolidated sand extends from approximately 25 feet to 85 feet below grade.      Contamination present up to 45 feet below grade.	The site is located adjacent to the Sacramento River. Groundwater is present approximately 18 feet below grade and flow is strongly incluenced by the Sacramento River and flows to the east.	TPH BTEX PAHs	Excavation and disposal;  Pump and treat;  In stu stabilization.	<i>In situ</i> stabilization implemented late 2012
Former Red Bluff MGP Red Bluff, CA	http://www.pge.com/about/e nvironment/taking- responsibility/mgp/red- bluff.shtml	Subsurface soil consists of up between 3 and 28 feet of debris containing fill material underlain by a sily clay / clayey silt with interbedded sand, grave, and finergrained sediments.  Contamination present in the fill material.	The site is bound to the east by the Sacramento River. Groundwater is present between 4 and 39 feet below grade and is heavily influenced by river level. Groundwater flows either east, or west, depending on river stage.	TPH BTEX PAHs	Excavation and disposal of shallow source soils;  In situ stabalization of deeper source soils.	Remedial Action approved

#### Table 8-1 - Nationwide MGP Site Summary

Bremeton Gas Works Site Bremerton, Washington

MGP Site Name & Location	Reference	Geologic Conditions	Groundwater / Surfacewater	Contaminants of Concern	Remedial Actions	Cleanup Status
Georgia MGP	m/stuff/contentmgr/files/0/50	Subsurface soil consists of up to 22 feet of fill underlain by 15 feet of alluvium above weathered bedrock.  Contamination present to the bedrock.	The site is bounded to the west by the Chattahoochee River.	BTEX PAHs	In situ stabilization; Excavation and disposal; Groundwater barrier.	Remedial Action complete
Nyack MGP Site Nyack, NY	http://www.dec.ny.gov/docs/r emediation_hudson_pdf/rod 34404601.pdf	Subsurface soil consists of up to 13 feet of fill underlain by native slity sand and glacial till layers. Sandstone bedrock was encountered approximately 40 feet below grade. Contamination present to the bedrock.	The site is bound to the north by the Hudson River.  The bedrock is a productive aquifer with the groundwater flowing upward through the bedrock. Groundwater generally flows toward the Hudson River.  River sediments have been impacted.	BTEX PAHs	Excavation and disposal;  "In situ stabilization;  In situ chemical oxidation;  Dredging and disposal.	Upland solidification complete. Sediment removal scheduled to begin in 2013
Manitowoc Former MGP Site Manitowoc, WI	/cleanup/manitowoc/pdfs/m	Subsurface soil consists of 3-10 feet of miscellaneous sand/sit/clay fill material overlying glacial deposits of sind with varying amounts of gravel, silt, and clay. Unconsolidated materials extend to at least 40 feet below grand and bedrock is estimated to be approximately 48 to 50 feet below grade.      Contamination present up to 27 feet below grade.	The site is bound to the northwest by the Manitowoc River. Groundwater is present between 5 and 22 feet below grade and flows towards the Manitowoc River.  River sediments have been impacted.	BTEX PAHs Cyanide	Shallow excavation and disposal;  In situ stabilization;  Pump and treat (carbon);  In situ stabilization for sediments failed;  Dredging.	Pump and Treat O&M Sediment dredging scheduled to begin December 2013
Kinston MGP Site Kinston, NC	http://www.neuselibrary.org/	Subsurface soils consist of gravel fill undertain by a fine to medium grained sand layer with some gravel and clay up to 21 feet below grade. The sandy layer is undertain by a siltclay which extends up to 45 feet below grade, followed by a silty sand extending to 55 feet below grade.  Contamination present up to 23 feet below grade.	The Neuse River borders more than 50% of the Site including the north, west, and southwest boundaries. Groundwater flow is to the southwest, towards the Neuse River.  River sediments have been impacted.	BTEX PAHs Cyanide	<i>In situ</i> stabalization; Institutional controls.	Remedy selected, awaiting implementation

BTEX = benzene, tolouene, ethylbenzene, and xylenes

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

MGP = manufactured gas plant NAPL = non-aqueous phase liquid

O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

Bremeton Gas Works Site Bremerton, Washington

								Potent	tial Sou		MGP-R		Contan	ninants				
Duol	liminary Contaminants of Detantial Concern		Page	an far Insk	ion		Fee	dstocks	and	(se	ee Note		o Drenne	duata		Potential Human Health and Environmental Concerns		
Prei	liminary Contaminants of Potential Concern		Reas	on for Incl	ision			Fuels			WIGP	Proces	s Bypro	aucts	(S)		(see Note 2	)
Contaminant Group	Contaminant Benzene	× Potential MGP-Related Constituents	× Other Sources (see Note 3)	× Detected in Previous Sampling Efforts	X Detected Above Initial PRGs	Other EPA Contaminants	× Gasoline	Diesel fuel oil	Coal/coke briquettes	Ash, clinker, cinder, slag, soot, bricks	× Spent scrubber media (tarry wood chips)	× Tar (potentially as DNAPL)	× Light oil (potentially as LNAPL)	× Gas Liquor and Emulsion (tar-water mixture)	× Spent purifier media (iron oxide, tarry wood chips)	× Human Health Risk (Carcinogen)	× Other Human Health Risks (non-Carcinogen)	× Toxicity to Ecological Receptors
l	Toluene	X	X	X			X				x	X	X	X	X		X	X
	Ethylbenzene	Х	Х	Х	Х		Х				Х	Х	Х	Х	Х		Х	Х
ĺ	Xylenes	Х	X	X	Х		Х		-		Х	Х	Х	Х	Х		X	Х
ĺ	1,2,3-Trichlorobenzene 1,2,4-Trimethylbenzene	Х	х	X			х	<b>-</b>			х	х	х	Х	х		X	Х
l	1,3,5-Trimethylbenzene			X								<u> </u>	<u> </u>				X	
l	1,4-Dichlorobenzene			Х												Х	Х	Х
l	1,4-Difluorobenzene			X	V													V
l	1,2-Dichloroethane 2-butanone		Х	X	Х											Х	X	X
l	Acetone			X													X	X
ds	Carbon disulfide			Х													Х	Х
Volatile Organic Compounds	Carbon Tetrachloride		Х	X	Х											Х	X	X
Ĕ	Chlorobenzene-d5 Chloroform		х	X	Х											Х	X	X
8	cis-1,3-Dichloropropene		X	X	X											X	X	^
gani	cis-1,2-Dichloroethene			Х													Х	
ō	Cyclochexane			Х													Х	
l iii	Isopropylbenzene			X													X	Х
No.	Methyl acetate Methylcyclohexane			X													Х	
1	Methylene chloride			X												Х	Х	Х
l	n-Butylbenzene			X													X	
l	n-Hexane			Х	Х												Х	Х
l	n-Propylbenzene			Х													Х	
l	Pentafluorobenzene p-Isopropyltoluene			X														
l	sec-Butylbenzene			X													Х	
l	Styrene			X													Х	Х
l	Tetrachloroethene			Х												Х	Х	Х
l	trans-1,3-Dichloropropene		X	X	X											X	X	.,
l	Trichloroethene Trichlorofluoromethane		Х	X	Х											Х	X	X
	Benzo(a)anthracene	Х	Х	X	Х				х	Х	х	Х		Х	Х	Х	X	X
	Benzo(b)fluoranthene	Х	X	X	Х				Х	Х	Х	Х		Х	Х	X	X	Х
	Benzo(k)fluoranthene	Х	Х	Х	Х				Х	Х	Х	Х		Х	Х	Х	Х	Х
s	Benzo(a)pyrene	X	X	X	X				X	X	X	X		X	X	X	X	X
Polycyclic Aromatic Hydrocarbons	Chrysene Dibenz(a,h)anthracene	X	X	X	X			-	X	X	X	X	-	X	X	X	X	X
Scar	Indeno(1,2,3-cd)pyrene	X	X	X	X				X	X	X	X	$\vdash$	X	X	X	X	X
ydrc	Acenaphthene	Х	X	X	X			Х	X	X	X	X		X	Х		X	X
Ę.	Acenaphthylene	Х	Х	Х	Х			Х	Х	Х	Х	Х		Х	Х		Х	Х
nat	Anthracene	X	X	X	X		-	X	X	X	X	X	<u> </u>	X	X		X	X
Aroi	Benzo(g,h,i)perylene Dibenzofuran	Х	Х	X	X			Х	Х	Х	Х	Х		Х	Х		X	X
i i	Fluoranthene	Х	х	X	X		<del>                                     </del>	х	х	Х	х	Х	$\vdash$	Х	Х		X	X
χ̈́ς	Fluorene	X	x	X	X			X	X	X	X	X		X	X		X	X
Po	Phenanthrene	Х	Х	Х	Х			Х	Х	Х	Х	Х		Х	Х		Х	Х
	Pyrene	X	Х	X	X			Х	X	X	Х	Х		Х	Х		X	X
	Methylpaphthalene, 1-	X	X X	X	X		X	X	X	X	X	X	X	X	X		X	X
	Methylnaphthalene, 2- Naphthalene	X	X	X	X		X	X	X	X	X	X	Х	X	X			Х

Bremeton Gas Works Site Bremerton, Washington

	Potential Sources of MGP-Related Contaminants (see Note 1)				Potential Human Health and													
Prel	liminary Contaminants of Potential Concern		Reas	on for Inclu	ısion		Fee	dstocks Fuels	and		MGP	Proces	s Bypro	ducts			nmental Co (see Note 2	
Contaminant Group	Contaminant	Potential MGP-Related Constituents	Other Sources (see Note 3)	Detected in Previous Sampling Efforts	Detected Above Initial PRGs	Other EPA Contaminants	Gasoline	Diesel fuel oil	Coal/coke briquettes	Ash, clinker, cinder, slag, soot, bricks	Spent scrubber media (tarry wood chips)	Tar (potentially as DNAPL)	Light oil (potentially as LNAPL)	Gas Liquor and Emulsion (tar-water mixture)	Spent purifier media (iron oxide, tarry wood chips)	Human Health Risk (Carcinogen)	Other Human Health Risks (non-Carcinogen)	Toxicity to Ecological Receptors
spunoc	1.1'-Biphenyl 1.2.4-Trichlorobenzene 2.4-Dimethylphenol 4-Methylphenol			X X X												X	X X X	X X X
Other Semi-volatile Organic Compounds	Acetophenone Benzyl butyl phthalate Benzaldehyde			X X X												Х	X X X	Х
-volatile On	Bis(2-ethylhexyl) phthalate Caprolactam Carbazole creosols	X		X							X	X		X	X	X	X X X	X
Other Semi	Dibenzofuran Di-n-butyl phthalate Isophorone	X		X X							X	X		X	X	×	X X X	X X X
<u> </u>	Pentachlorophenol Phenol Aluminum Antimony	х	Х	X X X	X						Х	Х		Х	Х	Х	X X X	X X X
	Arsenic Barium Beryllium		Х	X X X	X											Х	X X X	X X X
	Cadmium Chromium Cobalt		X	X X X	X X X												X X X	X X X
Metals <sup>4</sup>	Copper Iron Lead		X	X X X	X												X X X	X X X
	Manganese Mercury Nickel Selenium		X	X X X	X												X X X	X X X
	Silver Thallium Vanadium			X X X	X X												X X X	X X X
Polych Pestici	Zinc Norinated Biphenyls (PCBs) <sup>5</sup>		Х	X	X	X										X	X X	X X
Other	Cyanide, WAD Cyanide, total Sulfide	X X X				Х									X	X	X X X	X X X

#### Note

- 1) Contaminants of Potential Concern (COPCs) associated with MGP sources based on typical composition of MGP-related feedstocks and byproducts (see Section 2.3.1.1).
- 2) Potential Human Health and Environmental Concerns identified based on whether risk-based screening levels or potential ARARs for human health (carinogenic health effects), human health (non-carcinogenic health effects), or ecological health effects were identified during development of initial Preliminary Remediation Goals (PRGs) (see Section 6).
- 3) Other Sources include other historical operations at the site or regional sources of contamination.
- 4) Although previously detected at the Site, non-toxic metals (calcium, magnesium, sodium and potassium) are not included herein. Initial PRGs were not developed for these metals because they are essential nutrients that can be tolerated in high doses by living systems.
- 5) PCBs were previously analyzed for and not detected above reporting limits in soil or groundwater at the Site. However, the full standard list of PCB aroclors are COPCs for further evaluation.
- 6) The full standard list of pesticides, identified and quantified by EPA Method 8081B, are preliminary COPCs.

This table is not intended to be an exhaustive and complete preliminary list of Site COPCs. The RI/FS will include analysis of samples for the full standard list of analytes under each contaminant group. This list will be evaluated and revised as data is collected and specific contaminants can either be eliminated from the COPC list or are identified as Site COPCs.

### **Table 8-3 - Remedial Technologies for NAPL**

Bremerton Gas Works Site Bremerton, Washington

Dee Restrictions	NAPL General Response Actions	Remedial Technology	Process Options	Description
Institutional Controls  Use Restrictors  We Restrictors  Deed restrictions or encounts in exposure of engineered controls  Deed restrictions are of engineered controls  Deed restrictions are of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the search of the exposure of the exposure of the search of the exposure o			signs to control Site	Signs, fences, or other measures to prevent access to the Site.
Deed restrictions addressing sail disturbance analyze graunidators weeks	Institutional Controls	Use Restrictions	monitoring to prevent disturbance of	
Situr Wall   Situry Wall   Situry Wall   Situry Wall   Situry   Or in 38th missing of Bentonite with native soils.			addressing soil disturbance and/or	
Situ Treatment   Vertical basirers   Sheet Piles Vall   Water Injection   Control lateral movement of NAPL by pressure injecting hydraulic cements, clays, bentonite, and silicates into the formation through tighty speed borings using jetting tools.   A variety of heating methods, heating to temperature lifeting   Thermal Conductive heating   Thermal Conductive			Slurry Wall	backfilling with a low-permeability material (e.g., bentonite
Grout Curtain   Comments, clays, bentonite, and silicates into the formation through tightly spaced borings using jetting tools.	<i>In Situ</i> Containment	Vertical Barriers	Sheet Pile Wall	I
Low-Temperature Thermal Treatment			Grout Curtain	, ,
Thermal Treatment   Heating   Thermal Conductive Heating			Hot Water Injection	boiling point of water, increasing the mobility and solubility of
Thermal Conductive Heating  Steam Injection  Mild-Temperature Thermal Freatment  In Situ Treatment  In Situ Treatment  In Situ Treatment  In Situ Treatment  Treatment  Thermal Conductive Heating  Treatment  High-Temperature Thermal Treatment  High-Temperature Thermal Treatment  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Treatment  Thermal Treatment  Thermal Treatment  Stabilization  Stabilization  Stabilization  Stabilization  Stabilization  Stabilization  Stabilization  Thermical Conductive Heating  Solidification/Stabilization  Chemical Treatment  Thermal Conductive Heating  Solidification/Stabilization  Stabilization  Chemical oxidation  Chemical oxidation  Thermal Conductive Heating Stabilized by the propertion of the propert		Thermal		pumping from wells, and contaminants are treated. Heating can
Mid-Temperature Thermal Treatment				conduction from vertical heated wells, or by electrical resistance
Mid-Temperature Thermal Treatment Thermal Treatment Thermal Conductive Heating Thermal Conductive Heating Temperature Thermal Conductive Heating Treatment Thermal Conductive Heating Treatment Thermal Conductive Heating Treatment Thermal Conductive Thermal Conductive Heating Treatment Thermal Conductive Heating Thermal Conductive Thermal Conductive Heating Thermal Conductive Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Solidification/ Stabilization Stabilization Stabilization Stabilization Thermal Conductive Heating Thermal Conductive Heating Solidification/ Stabilization Thermal Conductive Heating Solidification/ Stabilization Thermal Conductive Heating Solidification/ Stabilization Stabilization Stabilization Stabilization Stabilization Stabilization Chemical oxidation Stabilization Stab			Steam Injection	
Thermal Conductive Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Heating   Flectrical Resistance   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical Flectrical Period   Flectrical F		Thermal		extraction, contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed
Heating Heating Heating Heating Heating Heating Heating Heating Heating Temperature Thermal Treatment Thermal Treatment Thermal Treatment  Thermal Conductive Heating  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Chemical Treatment  Chemical Treatment  Chemical Treatment  Chemical oxidation  Chemical oxidation  Chemical oxidation Treatment  NAPL Pumping  Pumping of NAPL from wells and tenches  Surfactant Enhanced Recovery Excavation  Excavation  Excavation  Disposal  Pick Situ Treatment  Heating  Heating  Meating  Meating  Meating  Meating  Meating  Migh- Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Solidification/ Solidification/ Stabilization  Solidification/ Solidificatio				vertical heated wells, or by electrical resistance when voltage is
Thermal Treatment Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating Thermal Conductive Heating  Thermal Conductive Heating  Thermal Conductive Heating  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Chemical Oxidation involves the injection of chemical oxidants into the subsurface to react with an destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate.  Pumping of NAPL from wells and trenches  Surfactant Enhanced Recovery  Pumping of mobilized MAPL and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with recirculating system.  Excavation  Excavation  Thermal  Thermal  Co-Burning  Co-Burning  NAPL is removed by excavating soil containing NAPL.  Combustion of coal tar or tar contaminated soil with coal in utility boilers and cement kilns.  When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized.  Reuse of recovered product.  Disposal of recovered  Treatment Disposal of recovered  Treatment Disposal of recovered  Treatment Disposal of recovered  Treatment Disposal of recovered  Treatment of NAPL via incineration at a hazardous waste	<i>In Situ</i> Treatment	High.		The subsurface is heated to temperatures above the boiling point of water, volatilizing or destroying (by pyrolysis) volatile and semivolatile organic compounds. Contaminated vapors are collected
Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Solidification/ Stabilization  Chemical Chemical Treatment  Chemical oxidation  Chemical oxidation  Chemical oxidation  Chemical oxidation  Chemical oxidation  Chemical oxidation  Chemical oxidation  Chemical oxidation involves the injection of chemical oxidants into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate.  Pumping of NAPL from wells and trenches  Surfactants are injected near NAPL zones in groundwater to mobilize the NAPL, and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with recirculating system.  Excavation  Excavation  Excavation  Co-Burning  Co-Burning  NAPL is removed by excavating soil containing NAPL.  Combustion of coal tar or tar contaminated soil with coal in utility boilers and cement kilns.  When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized.  Recycling of recovered NAPL  Disposal of recovered  Treatment of NAPL via incineration at a hazardous waste		Thermal		pumping from wells, and contaminants are treated. Heating can be performed by thermal conduction from vertical heated wells, or by electrical resistance when voltage is applied between
Chemical Treatment   Chemical oxidation   Into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate.		Stabilization	•	solidify or immobilize contaminants. Potential amendments include polymers, pozzolans, and cement. Amendments can be mixed with soil <i>in situ</i> using large-diameter augers, soil mixers, or
Removal  Surfactant Enhanced Recovery Excavation  Excavation  Thermal  Disposal  Off-Site Management  NAPL Pumping wells and trenches  Surfactants are injected near NAPL zones in groundwater to mobilize the NAPL, and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with recirculating system.  Co-Burning  Co-Burning  NAPL is removed by excavating soil containing NAPL.  Combustion of coal tar or tar contaminated soil with coal in utility boilers and cement kilns.  When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized.  Recycling of recovered NAPL  Disposal of recovered  Treatment of NAPL via incineration at a hazardous waste			Chemical oxidation	into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide,
Removal  Surfactant Enhanced Recovery  Excavation  Excavation  Excavation  Co-Burning  Incineration  Disposal  Off-Site Management  Disposal  Surfactant Enhanced Recovery  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  Pumping of mobilized NAPL  NAPL  Pumping of mobilized NAPL  NAPL  Pumping of mobilized NAPL  NAPL  Combustion of coal tar or tar contaminated soil with coal in utility boilers and cement kilns.  When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized.  Reuse of recovered product.  NAPL  Treatment of NAPL via incineration at a hazardous waste		NAPL Pumping	· -	Pumping to remove NAPL that accumulates in a well or trench.
Co-Burning   Combustion of coal tar or tar contaminated soil with coal in utility boilers and cement kilns.   When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized.	Removal	Enhanced		mobilize the NAPL, and then the mobilized NAPL is extracted. May be applied with injection-withdrawal technique or with
Ex Situ Treatment  Thermal  Incineration  Incineration  When soil or sediment containing NAPL is heated to temperatures above 1,400°F, contaminants are directly oxidized.  Recycling of recovered NAPL  NAPL  Disposal  Disposal of recovered  Treatment of NAPL via incineration at a hazardous waste		Excavation		NAPL is removed by excavating soil containing NAPL.  Combustion of coal tar or tar contaminated soil with coal in utility
Disposal  Off-Site Management  NAPL  Disposal of recovered  NEeuse of recovered product.  Treatment of NAPL via incineration at a hazardous waste	Ex Situ Treatment	Thermal		When soil or sediment containing NAPL is heated to temperatures
Disposal  Management  Disposal of recovered  Treatment of NAPL via incineration at a hazardous waste	<u> </u>	Off-Site		
NAPL via incineration treatment facility.	Disposal	Management	Disposal of recovered	

BTEX = benzene, tolouene, ethylbenzene, and xylenes

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

NAPL = non-aqueous phase liquid

O&M = operation and maintenance PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

Soil General Response	Remedial Technology	Process Options	Description
	Access Restrictions	Fences and warning signs to control Site access	Signs, fences, or other measures to prevent access to the property.
Institutional Controls	Use Restrictions	Use restrictions and monitoring to prevent disturbance of engineered controls  Deed restrictions	Covenant placed on the property that limits or prohibits activities that may interfere with a cleanup action or result in exposure to hazardous substances.
		addressing soil disturbance	
		Permeable soil cover	Placing clean soil on the surface provides a barrier that prevents exposure to underlying soil but allows storm water to infiltrate.
<i>In Situ</i> Containment	Capping	Low-permeability cap	Low-permeability caps may be constructed of low-permeability soil such as clay or an engineered material such as asphalt or concrete.  This cap would not only prevent exposure to underlying soils, but would also minimize stormwater infiltration through potentially contaminated materials, thereby reducing mobility of contaminants located in the unsaturated soil zone. Engineered materials could also be used in areas requiring a durable surface, such as high-traffic areas.
		Impervious cap	Impervious caps may be constructed of low-permeability soil such as clay or an engineered material such as asphalt or concrete, overlain by an additional impermeable layer. This cap would not only prevent exposure to underlying soils, but would also prevent stormwater from infiltrating through potentially contaminated soils beneath the cap, thereby reducing mobility of contaminants located in the unsaturated soil zone. Often combined with barrier wall technology to fully encapsulate soils.
	Physical Removal and Treatment	Passive venting of soil vapors	Passive soil venting is a less aggressive version of soil vapor extraction that is usually applied to prevent contaminated soil vapors from migrating into buildings or crawl spaces. In passive venting, soil vapors beneath a building foundation are vented to the atmosphere either through atmospheric pressure changes or by applying a low vacuum with a ventilation fan. Vented vapors can be passed through activated carbon for treatment if necessary.
		Soil vapor extraction	Soil vapor extraction applies a vacuum to subsurface soil to volatilize contamination and extract soil vapor. Vapor stream is treated above ground to remove contamination before discharge.
		Hot Water Injection	The subsurface is heated to temperatures less than the boiling point of water, increasing the mobility and solubility of NAPL and NAPL
	Low-Temperature Thermal Treatment	Electrical Resistance Heating	constituents. Contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed by injecting steam in vertical wells, thermal conduction from vertical
In Situ		Thermal Conductive Heating	heated wells, or by electrical resistance when voltage is applied between subsurface electrodes.
Treatment		Steam Injection	The subsurface is heated to temperatures near the boiling point of water, volatilizing or destroying (by pyrolysis) volatile organic
	Mid-Temperature Thermal Treatment	Electrical Resistance Heating	compounds. Contaminated vapors are collected using soil vapor extraction, contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed by injecting
		Thermal Conductive Heating	steam in vertical wells, thermal conduction from vertical heated wells, or by electrical resistance when voltage is applied between subsurface electrodes.
	High-Temperature Thermal Treatment	Thermal Conductive Heating	The subsurface is heated to temperatures above the boiling point of water, volatilizing or destroying (by pyrolysis) volatile and semi-volatile organic compounds. Contaminated vapors are collected using soil vapor extraction, contaminated liquids are removed by pumping from wells, and contaminants are treated. Heating can be performed by thermal conduction from vertical heated wells, or by electrical resistance when voltage is applied between subsurface electrodes.
		Vitrification	Soil is heated via electrical current to temperatures greater than 2,400°F, destroying contaminants and fusing soil into a glassy matrix.

### Table 8-4 - Remedial Technologies for Soil

Bremerton Gas Works Site Bremerton, Washington

Soil General Response	Remedial Technology	Process Options	Description
	Stabilization	Solidification/ Stabilization	Soil or sediment is stabilized by adding amendments to solidify or immobilize contaminants. Potential amendments include polymers, pozzolans, and cement. Amendments can be mixed with soil <i>in situ</i> using large-diameter augers, soil mixers, or similar equipment.
<i>In Situ</i> Treatment	Chemical Treatment	Chemical oxidation	Chemical oxidation involves the injection of chemical oxidants into the subsurface to react with and destroy organic contaminants. Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate, which have been shown to destroy a wide range of contaminants in soil.
		Bioventing	Bioventing supplies oxygen to unsaturated soil to increase aerobic biodegradation rates and may be designed to increase the air exchange rate through the soil.
	Bioremediation	Amendment Injection	Biodegradation of contaminants by indigenous soil microbes can be enhanced by amending soil with nutrients, moisture, and oxygen (typically provided by injecting air or solutions into wells or trenches).
Removal	Excavation	Excavation	Excavators, backhoes, and other conventional earth moving equipment are the most common equipment used to remove contaminated soil from upland areas.
	Physical	Solidification/ Stabilization	Amendments are added to excavated soil or sediment to immobilize and/or bind contaminants within the stabilized product. Depending on the proportion of amending agents, the end product may take on the form of a quasi-soil/concrete material that could later be used as bulk fill.
		Co-Burning	Combustion of Manufactured Gas Plant residues, such as coal tar and tar contaminated soil, with coal in utility boilers and cement kilns.
	Thermal	Thermal desorption	Low-temperature thermal desorption involves heating soils or sediments to temperatures between 200°F and 600°F until volatile and semivolatile chemicals of concern (COCs) such as benzene and naphthalene evaporate. Exhaust gases produced by the process are typically combusted.
		Incineration	When soil is heated to temperatures above 1,400°F, contaminants are directly oxidized.
Ex Situ Treatment	Chemical/ Physical	Particle washing	In particle washing, soil is put in contact with an aqueous solution to remove contaminants from the soil particles. The suspension is often also used to separate fine particles from coarser particles, allowing beneficial use of the coarser fraction (if sufficiently clean) at the Site.
		Solvent extraction	Solvent extraction is a variant of soil washing in which an organic solvent (rather than an aqueous solution) is put in contact with the soil to remove contaminants.
		Landfarming	Microbial population potentially enhanced with nutrients, moisture, and bioaugmentation to treat contaminated soil on lined beds with tilling and irrigation.
	Bioremediation	Biopiles	Microbial population potentially enhanced with nutrients, moisture, aeration, and bioaugmentation to treat contaminated soil in stockpiles.
		Bioreactor	Microbial population potentially enhanced with nutrients, moisture, aeration, and bioaugmentation to treat contaminated soil in enclosed reactor vessels.
Reuse	Asphalt Batching	Cold-Mix Asphalt Batching	Encapsulation of contaminant by blending residues, wet aggregate and asphalt emulsion at ambient temperature.
		Hot-Mix Asphalt Batching	Encapsulation of contaminant by blending residues, wet aggregate and asphalt emulsion at high temperature.
Disposal	Confined On-Site Disposal	Confined On-site disposal	Excavated soils exceeding applicable cleanup standards could potentially be placed on site in a specially designed upland confined disposal facility (CDF). Depending on the leachability of confined materials, the CDF could potentially include a liner and a liquid collection system to prevent leachate from contaminating groundwater.
	Off-Site Landfill Disposal	Subtitle D (Solid Waste) Subtitle C	Contaminated soils from the Site may be transported to an off-site, permitted disposal facility. This disposal method provides for secure, long-term containment of hazardous and non-hazardous solid wastes.
	L	(Hazardous Waste)	-

#### Notes:

BTEX = benzene, tolouene, ethylbenzene, and xylenes

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

NAPL = non-aqueous phase liquid

O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

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Groundwater General Response Actions	Remedial Technology	Process Options	Description				
Institutional Controls	Deed Restrictions	Deed restrictions to preclude drinking water use  Deed restrictions addressing groundwater wells	Covenant placed on property that limits or prohibits activities that may interfere with a cleanup action or result in exposure to hazardous substances.				
Monitored Natural Attenuation	Monitored Natural Attenuation	Groundwater Monitoring	Provides monitoring to document the presence and effectiveness of natural processes in removing or containing Site chemicals of concern (COCs).				
		Slurry Wall	Control lateral movement of contaminated groundwater by installing impermeable vertical barriers. Vertical barriers can be constructed of a				
	Vertical Barriers	Sheet Pile Wall	variety of materials and installation techniques, including driving or vibrating steel sheet piling, excavation of a trench and backfilling with a				
		Grout Curtain	low-permeability material (e.g., bentonite slurry), in situ mixing of bentonite with native soils, or pressure injecting hydraulic cement ar bentonite.				
<i>In Situ</i> Containment	Pumping	Pumping from vertical wells or trenches	Migration of contaminants dissolved in groundwater can be controlled by pumping groundwater from vertical wells or trenches, creating a capture zone within which groundwater flows toward the capture point.				
	Stormwater Controls	Targeted Infiltration	A hydraulic barrier can be created by collecting and infiltrating stormwater and forming a local groundwater "mound."				
	Stormwater controls	Reduced Infiltration	Hydraulic controls can reduce localized infiltration and seepage of stormwater in impacted areas along the shoreline.				
	Permeable Reactive Barrier	Sorptive/Reactive Wall	A 40-foot-deep trench may be excavated in the uplands and filled with permeable material that sorbs dissolved-phase contaminants, facilitatin further biodegradation and limiting contaminant migration toward marine sediment and surface water and offshore groundwater. A shallo trench could also excavated on the beach near the shoreline, but would be impacted by brackish water and tidally-influenced groundwater gradients.				
<i>In Situ</i> Treatment	Chemical Treatment	Chemical Oxidation	Chemical oxidation involves the injection of oxidant solutions into saturated groundwater to react with and destroy organic contaminants.  Common oxidants include hydrogen peroxide, potassium permanganate, ozone, and sodium persulfate.				
		Amendment Injection	Injecting compounds, such as peroxides, oxygen-releasing compound, or nutrients, that enhance degradation of contaminants.				
	Bioremediation	Biosparging	Biosparging involves the injection of oxygen, and sometimes nutrients, to groundwater to enhance aerobic bioattenuation of organic compounds. For volatile contaminants, soil vapor extraction or bioventing may be concurrently applied for unsaturated soil.				
Removal	Groundwater Extraction	Pumping from Vertical Wells or Trenches	Groundwater can be removed from the subsurface by pumping fluids from wells or trenches.				
		Adsorption	Granular activated carbon (GAC) can be used to remove organic contaminants. Contaminated groundwater is passed through a bed of GAC, and hydrophobic organic compounds in solution adsorb onto the carbon until the carbon becomes depleted or saturated. Depleted GAC may be regenerated or disposed off Site.				
<i>Ex Situ</i> Treatment	Physical/ Chemical	Air Stripping	Contaminated groundwater and air are typically passed counter-current through a tower, and volatile contaminants (such as benzene and, to a lesser extent, naphthalene) transfer from the water to the air. The contaminant-laden air is usually treated by activated carbon and then discharged to the atmosphere.				
		Advanced Oxidation Processes	Involves adding chemicals that directly oxidize organic contaminants in water. Process options include ozonation, hydrogen peroxide (with or without catalysts such as Fenton's Reagent or ultraviolet light), and permanganate.				
	Biological	Biotreatment	Contaminated groundwater is passed through a biological reactor in which a contaminant-degrading microbial culture is maintained, generally by adding nutrients and oxygen and controlling temperature, pH, and other parameters. Process options include bioslurry reactors, fixed-film bioreactors, and constructed wetlands.				
	Off-Site Management	Discharge to Sanitary Sewer	Groundwater is discharged to the local sanitary sewer system. Pre- treatment of groundwater may not be required if concentrations of chemicals of concern (COCs) meet discharge criteria. Water containing high concentrations of solids (e.g., from construction dewatering) would likely need to be passed through a settling tank or filter to meet discharge requirements.				
Disposal		Discharge to Surface Water	Extracted groundwater may also be discharged to surface water, although this discharge option would likely require a National Pollutant Discharge Elimination System (NPDES) permit. Water discharged to surface water would have to meet strict water quality requirements and would likely require treatment before discharge.				
	On-Site Management	Re-introduction to Groundwater	Extracted groundwater may also be discharged on site to groundwater via infiltration galleries or injection wells. Contaminated groundwater would likely require treatment before discharge via this method.				

BTEX = benzene, tolouene, ethylbenzene, and xylenes

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

NAPL = non-aqueous phase liquid O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

## Table 8-6 - Remedial Technologies for Sediment Bremerton Gas Works Site

Sediment General Response Actions	Remedial Technology	Process Options	Description
		Governmental advisories and public outreach on fish/shellfish consumption	
Institutional Controls	Use Restrictions	Easements or restrictive covenants to limit activities which may damage the remedy or increase the potential for exposure	Institutional controls are measures undertaken to limit or prohibit activities that may interfere with a cleanup action or result in exposure to hazardous substances.
		Monitoring and notification of waterway users to restrict specific activities to protect the remedy	
	Monitored Natural Recovery	Monitored Natural Recovery	A passive remedial approach which relies on monitoring of ongoing, natural processes (physical, biological, and/or chemical mechanisms) that act together to reduce the risk (bioavailability and/or toxicity) of the Site COCs. Monitoring is required to evaluate the effectiveness and frequently includes multiple lines of evidence.
Monitored Natural Recovery	Enhanced Natural Recovery	Thin-Layer Sand Placement	Thin-layer placement normally accelerates natural recovery by adding a layer of clean sediment over contaminated sediment. The acceleration can occur through several processes, including increased dilution through bioturbation of clean sediment mixed with underlying contaminants. Thin-layer placement is typically different than the <i>in situ</i> isolation caps, because it is not designed to provide long-term isolation of contaminants from benthic organisms.
<i>In Situ</i> Containment	Capping (Non- reactive)	Engineered Sand Cap	An engineered sand cap consists of a layer of granular material placed over contaminated sediments to contain and isolate them from the biologically active surface zone. Engineered caps may also include erosion protection or stability layers such as geosynthetics or armoring materials.
		Post-Dredge Residuals Management Layer	Similar to cap placement methods described above, with the exception that granular material is applied after dredging to manage residual contamination resulting from dredging. In some cases, a reactive media may be included in the residuals/backfill layer.
In Situ	Physical/ Chemical	Permeable Reactive Cap	A permeable reactive cap includes a reactive material (such as organoclay, coke, coal, or activated carbon) and similar to a sand cap is placed over contaminated sediments to isolate and contain the contaminated sediments. The reactive material also provides treatment by sorping or binding COCs (dissolved and/or NAPL) and further limiting migration into overlying sediment porewater and surface water.
Treatment		Stabilization	This technology involves adding amendments to in situ sediment that immobilize and/or bind contaminants within the stabilized media.
	Bioremediation	Amendment Injection	Biodegradation of contaminants by indigenous soil microbes can be enhanced by amending soil with nutrients, moisture, and oxygen (typically provided by injecting into wells or trenches).

### **Table 8-6 - Remedial Technologies for Sediment**

Bremerton Gas Works Site Bremerton, Washington

Sediment General Response Actions	Remedial Technology	Process Options	Description			
Removal	Dredging	Hydraulic	Dredging is the removal of sediment in the wet and is primarily accomplished with hydraulic or mechanical equipment. Hydraulic dredging removes and transports sediment with entrained water in a slurry. Mechanical dredging uses mechanical equipment/force to dislodge and excavate sediment in the wet. Dredging effectiveness may be limited by resuspension, release			
nemova	Dieuging	Mechanical	of COCs (i.e., dissolved, particles, and sheens) to water and volatilization to air during dredging, and residual COCs remaining after dredging (USACE 2008). These effects may be reduced by use of containment (e.g., sheet pile, silt curtains) and best management practices.			
	Physical	Physical Separation	The volume of excavated or dredged contaminated materials may be reduced by physically separating the materials into two or more fractions that can be handled separately.			
Ex Situ	,	Stabilization	This technology involves adding amendments to excavated sediment that immobilize and/or bind contaminants within the stabilized media.			
Treatment	Thermal	Thermal Desorption	Low-temperature thermal desorption involves heating soils or sediments to temperatures between 200°F and 600°F until volatile and semivolatile COCs such as benzene and naphthalene evaporate. Exhaust gases produced by the process are typically combusted.			
		Incineration	When sediment is heated to temperatures above 1,400°F, contaminants are directly oxidized.			
	On-Site Beneficial	Sand/Aggregate Reclamation	Dredged material with high sand contents that undergo particle separation may be available for use as concrete aggregate or general upland fill.			
	Use	Topsoil Feedstock	Dredged material may be used as non-organic feedstock for topsoil (i.e., material would be blended with organics).			
	Confined On-Site	Confined On-site Disposal	Removed sediments exceeding applicable cleanup standards could potentially be placed on Site in a specially designed upland CDF. Depending on the leachability of confined materials, the CDF could potentially include a liner and a liquid collection system to prevent leachate from contaminating groundwater.			
Disposal	Disposal	Near-shore Confined Disposal Facility (CDF)	Removed sediments exceeding applicable cleanup standards could potentially be placed on Site in a specially designed CDF built along the shoreline. Construction would require significant filling and conversion of aquatic lands.			
		Contained Aquatic Disposal (CAD)	Dredged sediments may be consolidated and disposed of in a deep aquatic excavation adjacent to the Site and capped with clean material.			
	Off-Site Landfill	Subtitle D (Solid Waste)	Contaminated sediments from the Site may be transported to an off-Site, permitted disposal facility. This disposal method			
	Disposal	Subtitle C (Hazardous Waste)	provides for secure, long-term containment of hazardous and non-hazardous solid wastes.			

#### Notes:

 ${\sf BTEX} = {\sf benzene}, \, {\sf tolouene}, \, {\sf ethylbenzene}, \, {\sf and} \, \, {\sf xylenes}$ 

COCs = chemicals of concern

cPAHs = carcinogenic polycyclic aromatic hydrocarbons

NAPL = non-aqueous phase liquid

O&M = operation and maintenance

PAHs = polycyclic aromatic hydrocarbons

TPH = total petroleum hydrocarbons

U.S. Army Corps of Engineers (USACE), 2008, Technical Guidelines for Environmental Dredging of Contaminated

References: Sediments, ERDC/EL TR-08-29, September 2008.

### Table 9-1 - Summary of Existing Information and Data Gaps - Uplands

Bremerton Gas Works Site Bremerton, Washington

Remdial Investigation/Feasbility Study Information Needs by	Existing Information	Data Gaps	Recommended Data Collection
Торіс			
Physical Characteristics			
Characteristics of water-bearing zones	Soil stratigraphy and observed/measured groundwater occurrence from previous investigations identifies a water-bearing zone in clean to silty glacial sands at depths of 15 to 41 feet below surface.	bearing zones and aquitards.	<ul> <li>Soil borings to evaluate soil stratigraphy and identify waterbearing zones and aquitards.</li> <li>Soil samples from borings for laboratory measurement of physical parameters that may include grain size, porosity, bulk density, and total/fraction organic carbon.</li> <li>Slug tests at select site wells to measure hydraulic conductivity in each saturated stratigraphic horizon and in different water-bearing zones (if applicable).</li> </ul>
Groundwater flow direction and gradient	Manual groundwater level measurements collected at eight wells in 2007 were used to evaluate groundwater flow direction and gradient.	<ul> <li>Groundwater flow direction and horizontal/vertical gradients.</li> <li>Seasonal variability in water levels and groundwater gradients.</li> <li>Influence of precipitation/surface water infiltration on groundwater levels.</li> <li>Influence of tidal fluctuation on groundwater levels.</li> </ul>	<ul> <li>Continuous water levels at site wells and in the Narrows using pressure transducers.</li> <li>Precipitation amounts recorded at area weather stations.</li> </ul>
Groundwater geochemistry	None.	Location of salt water intrusion and extent of groundwater- surface water interaction.	<ul> <li>Groundwater samples will be collected from site wells for field measurements and laboratory analysis of conventional geochemical parameters, salinity.</li> </ul>
Nature and Extent of Contamination			
Identify and evaluate source areas	Historical review of Gas Works operations identifies potential source areas.	<ul> <li>Identified potential source areas have not been sufficiently investigated.</li> <li>Potential locations of some potential sources (e.g., tar pits, transfer piping) are unknown or roughly estimated.</li> </ul>	<ul> <li>Ground-penetrating radar to identify potential subsurface features.</li> <li>Advance soil borings and/or complete test pits in and around potential source areas, including former process and residuals management areas, including the tar pit, residue cistern, tar wells, and in the ravine fill area.</li> <li>Visually observe and record soil stratigraphy and indications of contamination.</li> </ul>
Evaluate COPCs to determine COCs	Surface and subsurface soil and groundwater samples collected in 2007 and 2008 were analyzed for metals, petroleum hydrocarbons, SVOCs, VOCs and PCBs.	Presence of COPCs previously not evaluated (e.g., cyanide).	Soil and groundwater samples will be collected for chemical analysis of COPCs to refine COC list.
Define nature and extent of COCs in soil	Soil samples collected in 2007 and 2008 identified concentrations of metals, PAHs, and VOCs exceeding PRGs.	<ul> <li>Current nature and extent of COCs in soil.</li> <li>Presence, nature, and extent of COPCs previously not evaluated.</li> </ul>	Soil samples will be collected from soil borings and test pits in source areas and surrounding the Site to establish horizontal and vertical limits to the extent of comtamination. Soils will be submitted for chemical analysis of COCs.

## Table 9-1 - Summary of Existing Information and Data Gaps - Uplands

Bremerton Gas Works Site Bremerton, Washington

Remdial Investigation/Feasbility Study Information Needs by	Existing Information	Data Gaps	Recommended Data Collection
Торіс			
Define nature and extent of COCs in groundwater	Groundwater samples collected in 2007 and 2008 identified concentrations of metals, SVOCs, and VOCs exceeding PRGs.	<ul> <li>Current nature and extent of COCs in groundwater.</li> <li>Seasonal variability of COCs in groundwater.</li> </ul>	<ul> <li>Groundwater samples may be collected from soil borings if encountered to evaluate presence of COCs and inform well placement.</li> <li>Install monitoring wells to evaluate impacts in source areas and establish horizontal and vertical limits to the extent of contamination. Groundwater samples will be collected from monitoring wells for chemical analysis of COCs.</li> </ul>
Define nature and extent of NAPL	Previous investigations have indicated that NAPL may be present.	<ul> <li>Presence/absence of NAPL.</li> <li>Chemical composition of NAPL.</li> <li>Lateral and vertical boundaries of NAPL occurences.</li> </ul>	<ul> <li>Advance soil borings and/or complete test pits in former Gas Works operations and residuals management areas, including the tar pit, residue cistern, tar wells, and in the ravine fill area. Visually observe and record soil stratigraphy and NAPL occurrences.</li> <li>Include monitoring wells screened appropriately to monitor LNAPL (across water table) and DNAPL (above aquitards). Monitor wells for LNAPL and DNAPL presence.</li> <li>Submit representative soil samples and/or NAPL collected from soil borings, test pits, or wells for chemical analysis to characterize NAPL chemistry.</li> <li>If NAPL is identified to be present: advance additional soil borings for deeper NAPL occurences and test pits for shallow NAPL occurences in areas requiring more precise definition of NAPL occurrences.</li> </ul>
Evaluate potential for recontamination from other area sites  Contaminant Fate and Transport	Soil and groundwater samples that have been collected from borings and wells located upgradient of the Gas Works property show potential impacts in groundwater south of the property. Limited available data do not show impacts from bulk fuel facilities east of Pennsylvania Avenue or west of Thompson Drive extending onto the Gas Works property.	Potential impact from adjacent bulk fuel facilities and upgradient industrial sites.	Soil and groundwater data collected from soil borings, test pits, and monitoring wells upgradient of the former Gas Works property will be compared to evaluate the extent of contaminants exceeding screening criteria that are associated with the Gas Works site and potential contributions from other area contaminant sources.
NAPL migration pathways	NAPL may be present in the subsurface. MGP-related products	Nature and extent of NAPL (see above)	Characterize soil characteristics, NAPL characteristics, and
2 mgradon patriways	include both LNAPL and DNAPL.	NAPL mobility, including NAPL physical characteristics and soil lithology/physical properties	
Soil-to-groundwater pathway	Concentrations of Gas Works-associated constituents have been detected above soil and groundwater PRGs.	Leaching potential from contaminated soils.	<ul> <li>Include TOC in soil testing program.</li> <li>Collect groundwater chemistry data along groundwater flowpaths.</li> </ul>

#### Table 9-1 - Summary of Existing Information and Data Gaps - Uplands

Bremerton Gas Works Site Bremerton, Washington

Remdial Investigation/Feasbility Study Information Needs by	Existing Information	Data Gaps	Recommended Data Collection
Topic			
Soil-to-surface water pathway	Concentrations of Gas Works-associated constituents have been detected above soil PRGs.	Discharge of contamination through stormwater runoff.	Characterize contamination in exposed surface soil, catch basins sediments, and surface water discharging at outfalls.
Groundwater-to-surface water pathway	Concentrations of Gas Works-associated constituents have been detected in groundwater above surface water PRGs.	<ul> <li>Groundwater transport parameters (velocity, pathway).</li> <li>Attenuation parameters.</li> </ul>	<ul> <li>Include natural attenuation parameters in groundwater testing program.</li> <li>Characterize hydrogeology and chemical nature and extent (see above). Data may be incorporated into hydrogeologic and fate and transport models.</li> <li>Groundwater monitoring program to assess seasonal variability and long-term trends.</li> </ul>
Soil-to-air and groundwater-to-air pathway	Concentrations of Gas Works-associated constituents have been detected above current soil and groundwater PRGs.	Potential impacts to future indoor air.	Soil and groundwater data to be used with vapor transport modeling.
Human Health and Ecological Risk Assessment			
Assess potential receptors and exposure pathways		<ul> <li>Potential risk to human health through direct contact with soil, ingestion of groundwater, and inhalation via vapor intrusion.</li> <li>Potential risk to ecological receptors through direct contact with soil.</li> </ul>	Soil and groundwater chemical analytical results will be compared to human health and ecological risk-based criteria.

#### Notes:

BTEX = benzene, toluene, ethylbenzene and xylenes

COC = chemical of concern

COPC = chemical of potential concern

Cs-137 = Cesium 137 isotope

CSL = Cleanup Screening Level

CSO = combined sewer overflow

DNAPL = dense non-aqueous phas liquid

LNAPL = light non-aqueous phase liquide

MGP = manufactured gas plant

NAPL = non-aqueous phase liquid

NOAA: National Oceanic and Atmospheric Administration

PAHs = Polycyclic aromatic hydrocarbons

PCB = polychlorinated biphenyl

PRG = preliminary remediation goal

SCO = Sediment Cleanup Objective

SMS = Washington Sediment Management Standards regulations (WAC-173-204)

SVOC = semivolitile organic compound

TOC = total organic carbon

VOC = volatile organic compound

### Table 9-2 - Summary of Existing Information and Data Gaps - Sediments

Bremerton Gas Works Site Bremerton, Washington

Remedial Investigation/Feasibility Study Information Needs	Existing Information	Data Gaps	Recommended Data Collection
by Topic			
Nature and Extent of Contamination			
Assess presence of chemical contaminants associated with historical Gas Works operations.	<ul> <li>Gas Works operational history is well documented.</li> <li>MGP-associated contaminants typically include PAH compounds, selected VOCs (i.e., BTEX compounds), cyanide and dibenzofuran.</li> <li>Surface sediment PAH concentrations within the intertidal beach areas have been extensively sampled.</li> <li>Some testing for other parameters (SVOCs, metals, and VOCs) has also been performed on a more limited basis.</li> </ul>	<ul> <li>Sampling has not been performed in areas offshore of the former Gas Works dock.</li> <li>Testing has not been performed for cyanide in sediments.</li> <li>Testing for alkylated PAHs has not been performed (these parameters are useful in discriminating PAH sources in sediments).</li> </ul>	<ul> <li>Collect surface sediment samples from Gas Works dock area.</li> <li>Analyze sediment samples in selected areas for cyanide.</li> <li>Analyze sediments samples in selected areas for alkylated.</li> <li>PAH to document the "fingerprint" of MGP-associated PAH.</li> </ul>
Identify chemical contaminants potentially associated with other historical activities within the Site.	<ul> <li>Other potentially significant uses of the Site and vicinity include ravine fill, oil handling, CSO/stormwater discharges, adjacent marina operations and miscellaneous industrial operations on the Sesko and McConkey properties.</li> <li>Some testing for other parameters besides PAH compounds (semivolatiles, metals and VOCs) has been performed on a limited basis.</li> </ul>	<ul> <li>Sampling near non-MGP sources is not sufficient to finalize list of site-associated contaminants.</li> <li>Testing has not yet been performed offshore of former Sesko Oil dock.</li> <li>Testing for alkylated PAHs has not been performed (these parameters are useful in discriminating PAH sources in sediments).</li> </ul>	<ul> <li>Collect surface sediment samples from former Sesko dock area.</li> <li>Analyze sediment samples in selected areas for additional parameters to finalize list of site-associated COCs.</li> <li>Analyze sediment samples in selected areas for alkylated PAH to evaluate "fingerprint" and potential presence of non-MGP sources within the Site.</li> </ul>
Define the lateral extent of Site-associated COCs in surface sediment, including the boundary between Site-associated contamination, and contamination from other inputs.	<ul> <li>Surface sediment PAH concentrations within the intertidal beach areas have been extensively sampled.</li> <li>Some testing for other parameters (semivolatiles, metals and VOCs) has also been performed on a limited basis.</li> <li>Extensive data are available documenting sediment quality within Port Washington Narrows and Dyes Inlet. Those data indicate elevated PAH concentrations and the presence of certain other contaminants.</li> </ul>	<ul> <li>The lateral extent of site-associated PAH contamination has not been determined within Port Washington Narrows.</li> <li>Given the presence of elevated PAH concentrations in other sediments, additional sampling and "fingerprint" data will be needed to define the boundary between Site-associated PAH contamination and PAH contamination from other inputs.</li> <li>If other site-associated COCs are confirmed, then the lateral extent of these COCs in surface sediments will need to be determined, including the boundary between Site-associated contamination and contamination from other inputs.</li> </ul>	<ul> <li>Collect surface sediment samples from across the initial study area and analyze for selected parameters.</li> <li>Conduct surface sediment samples at selected locations outside the initial study area to evaluate other influences on sediment quality and the boundary between site-associated and other contaminant sources.</li> </ul>
Define the vertical extent of Site-associated COCs in subsurface sediment, including the potential presence of subsurface hydrocarbon deposits (i.e., sheen or NAPL).	Subsurface testing has been performed in the western portion of the intertidal beach to evaluate the vertical extent of PAH contamination and hydrocarbon sheen in that area. Results demonstrated that sediment contamination levels decreased rapidly (i.e., within a few feet) with depth, and the area containing subsurface hydrocarbon sheen was very limited.	<ul> <li>Subsurface testing has not been performed in other areas of the beach. The depth of contamination is therefore not defined in those areas.</li> <li>No surface or subsurface testing has been performed areas offshore of the former MGP dock.</li> <li>Core sampling data are not yet sufficient to assess whether subsurface hydrocarbon deposits (sheen or NAPL) may be present in subsurface sediments other than in the western beach area.</li> </ul>	<ul> <li>Conduct sediment core sampling and chemical analysis within portions of the initial study area to assess the vertical extent of PAH contamination.</li> <li>Include sufficient core sampling locations in nearshore and offshore areas to assess the potential presence of susurface hydrocarbon deposits (sheen or NAPL).</li> </ul>

### Table 9-2 - Summary of Existing Information and Data Gaps - Sediments

Bremerton Gas Works Site Bremerton, Washington

Remedial Investigation/Feasibility Study Information Needs by Topic	Existing Information	Data Gaps	Recommended Data Collection
Human Health & Ecological Risk Assessment			
Assess the site-specific partitioning behavior of PAHs in sediments.	Literature data can be used to estimate potential partitioning of PAH compounds between sediment and porewater. However, these methods may not capture site-specific factors.		<ul> <li>Conduct paired analysis of bulk sediment and porewater PAH concentrations in selected study areas for analysis of site- specific partitioning behavior.</li> </ul>
Assess potential impacts of site-associated COCs to benthic receptors.	<ul> <li>The potential for benthic impacts can be assessed using bulk sediment chemistry (to be defined as described above) along with toxicity threshold values such as the SMS SCO and CSL values, and/or the EPA narcosis toxicity model.</li> <li>Porewater PAH data may be used directly to assess potential benthic toxicity using the EPA narcosis toxicity model.</li> </ul>	<ul> <li>Site-specific bioassay testing could be used along-side bulk sediment chemistry and porewater testing data to assess potential benthic impacts.</li> <li>The need for bioassay testing can be assessed after review of bulk sediment chemistry and porewater PAH data to be collected as described above.</li> </ul>	Contingent Activity: If applicable, based on review of bulk sediment chemistry and porewater testing data, collect sediment samples from selected areas for confirmational bioassay testing. This testing could be used to verify predicted impacts and refine the lateral extent of those impacts.
Assess potential for site-associated sediment contaminants to accumulate in the tissues of aquatic organisms.	Literature data can be used to estimate potential uptake of PAH or other contaminants in the tissues of aquatic organisms. Reliance on literature data may not capture site-specific factors.		<ul> <li>Develop estimates of tissue concentrations based on bulk sediment and porewater testing data and literature-based biota-sediment accmulation factors.</li> <li>Contingent Activity: If warranted, use tissue testing (preferred) or laboratory bioaccumulation testing (alternate) to directly assess the potential accumulation of site-associated COCs in selected aquatic organisms.</li> </ul>
Document the types and quantities of aquatic species present in the vicinity of the Site and potentially relevant to human health and/or ecological risk evaluations.	<ul> <li>Previous habitat and fish/shellfish resource surveys have been performed in the Port Washington Narrows and Dyes inlet areas, documenting locally-abundant fish and shellfish species.</li> <li>Information regarding current and proposed shellfish growing areas, and historical patterns of fishing and shellfish harvesting are available through state and tribal agencies.</li> <li>Patterns of tribal seafood consumption have been identified in previous surveys of the Suquamish, Tulalip and Squaxin nations.</li> </ul>	<ul> <li>Additional information is required to document the habitat conditions and the types of seafood species present within Port Washington Narrows near the Site.</li> <li>The sustainable shellfish yield for the Site has not been defined. Such information will be helpful in applying shellfish consumption rates documented in the EPA Region 10 Tribal Framework for Selecting Fish and Shellfish Consumption Rates to the baseline risk assessment.</li> </ul>	<ul> <li>Conduct surveys of aquatic habitat and fish/shellfish resources at and near the Site within Port Washington Narrows.</li> <li>Define the potential shellfish yield for the Site based on surveys of similar properites within the Port Washington Narrows area.</li> </ul>
Evaluate potential site-associated water quality impacts as necessary to support exposure assessments in the human health and ecological risk assessments.	No surface water data are currently available for the Site. Regional studies have documented anthropogenic surface water contaminant inputs to Port Washington Narrows and Dyes Inlet, including but not limited to stormwater and CSO discharges. Any Site-specific sampling of surface water quality will need to consider potential off-site sources for measured water quality parameters.	needs.	Analyze surface water samples for site-associated COCs.  Samples to be collected from both within the initial study area and at selected background stations within Port Washington Narrows east and west of the Site.

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#### Table 9-2 - Summary of Existing Information and Data Gaps - Sediments

Bremerton Gas Works Site Bremerton, Washington

Remedial Investigation/Feasibility Study Information Needs	Existing Information	Data Gaps	Recommended Data Collection
by Topic			
Sediment Stability and Recovery Processes			
Assess potential near-bottom currents on long-term sediment	Peak tidal currents within Port Washington Narrows are	Near-bottom tidal currents can be significantly different than	Conduct empirical measurements of near-bottom and mid-
stability within the Site and immediate vicinity.	understood from existing studies (e.g., NOAA tide and current	open-water, mid-channel currents due to local and edge	channel tidal currents for use in an analysis of sediment
	data).	effects. No near-bottom current data are available for the Site	stability.
	Sediment texture and particle size will be defined during	or vicinity.	
	surface sediment testing as described above.		
Quantify sedimentation rates using geochronology cores and	Geochronology studies have been performed in several areas	Sedimentation rates can vary with location. No	Contingent Activity: If warranted, quantify net sedimentation
radio-dating.	of Puget Sound, documenting a general pattern of	sedimentation rate data are available for Port Washington	rates near the Site using geochronology test methods (i.e., thin-
	sedimentation.	Narrows areas near the Site.	section cores analyzed with Cs-137 radio-dating).

Notes:

BTEX = benzene, toluene, ethylbenzene and xylenes

COC = chemical of concern

Cs-137 = Cesium 137 isotope

CSL = Cleanup Screening Level

CSO = combined sewer overflow

EPA = U.S. Environmental Protection Agency

MGP = manufactured gas plant

NAPL = Non-aqueous phase liquid

NOAA = National Oceanic and Atmospheric Administration

PAH = polycyclic aromatic hydrocarbons

SCO = Sediment Cleanup Objective

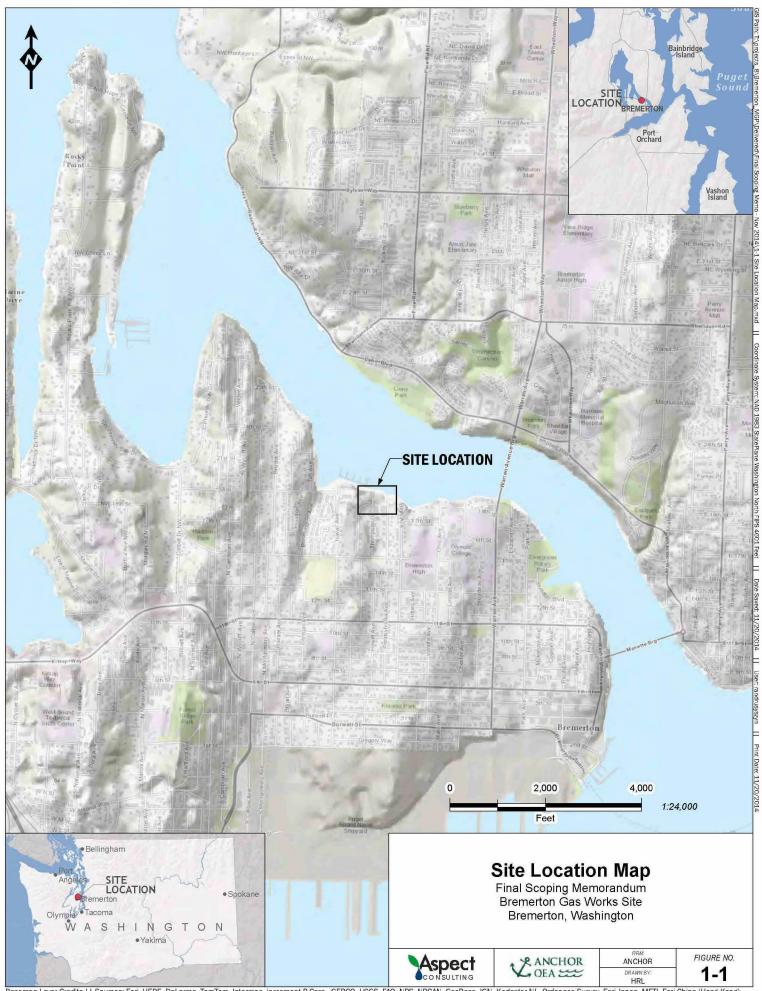
SMS = Washington State Sediment Management Standards regulations (WAC-173-204)

SVOC = semi-olatile organic compound

VOC = volatile organic compound

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# **FIGURES**



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